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Pursuant to 37 CFR 1.53(b), transmitted herewith for filing is the patent application of

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Title:

"METHODS AND COMPOSITIONS FOR THE TREATMENT AND PREVENTION OF TUMORS, TUMOR-RELATED DISORDERS AND CACHEXIA"

Priority Claim (35 U.S.C. 119) is made, based upon:

<u>Japanese Patent Application No.</u>	<u>Filing Date</u>
Hei 9-354499	December 24, 1997
Hei 10-15306	January 28, 1998
Hei 10-204907	July 21, 1998
Hei 10-269444	September 24, 1998

Enclosed herewith are:

- ☒ Specification (Description, Claims, Abstract): Pages 1 - 87; Number of claims 1 - 67
- ☒ Declaration and Power of Attorney (executed)
- ☐ **TO BE POST-FILED: SIGNED DECLARATION UNDER 37 CFR 1.53(f) UNEXECUTED DECLARATION ENCLOSED FOR INFORMATION PURPOSES ONLY**
- ☐ _____ Sheets of drawings, Figures 1 - _____ ☐ Formal ☐ Informal
- ☒ Assignment and Form PTO-1595 Recordation Form Cover Sheet and Check for \$40.00
- ☐ Certified copy (ies) of priority document(s) identified above
- ☐ Information Disclosure Statement including Form PTO-1449
- ☐ Preliminary Amendment
- ☐ Verified Statement(s) Claiming Small Entity Status
- ☒ Receipt Postcard

	Number Filed	Number Extra	Rate	Calculations
Total Claims	<u>67</u> - <u>20</u> = <u>47</u>		x \$18.00 =	\$ <u>846.00</u>
Independent Claims	<u>4</u> - <u>3</u> = <u>1</u>		x \$78.00 =	\$ <u>78.00</u>
MULTIPLE DEPENDENT CLAIMS			+ \$260.00 =	\$ <u>-</u>
			BASIC FEE	\$ <u>760.00</u>
			Total of above Calculations	\$ <u>1,684.00</u>

To the extent not tendered by check, authorization is given to charge any fees under 37 CFR 1.16 and 1.17 during pendency of the application, or to credit any overpayment, to Deposit Account No. 06-1378. Duplicate copy of this letter is enclosed.

FRISHAUF, HOLTZ, GOODMAN, LANGER & CHICK, P.C.

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HERBERT GOODMAN Reg. No. 17,081

**METHODS AND COMPOSITIONS FOR THE TREATMENT AND
PREVENTION OF TUMORS, TUMOR-RELATED DISORDERS AND
CACHEXIA**

Background to the Invention

5 The present invention relates to the use of certain compounds, specifically cyclooxygenase-2 inhibitors (hereinafter referred to as "COX-2 inhibitors") for the treatment and prevention of tumors and tumor-related disorders and cachexia.

10 Cachexia is a systemic disease of which the cardinal symptoms are progressive weight loss, anemia, edema, loss of appetite and so forth. It may occur as a side-effect of certain chronic diseases, such as malignant tumors, tuberculosis, diabetes, blood diseases, endocrine diseases, infections and acquired immune deficiency syndrome. When cachexia occurs as a result of the presence of a malignant tumor, even if the administration of antitumor drugs to the patient with malignant tumor is effective and antitumor effects are experienced, there is normally no improvement in
15 the cachexia because of adverse effects such as the myelotoxicity which may be caused by the antitumor drug.

 The treatment of cachexia is often very difficult for the following reasons:

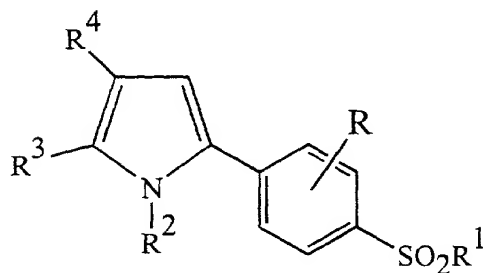
 Since the strength of a patient is greatly depleted as cachexia progresses, it may become impossible to continue treatment using antitumor drugs (which generally
20 exhibit a high level of toxicity), and this thereby becomes an obstacle to the treatment of the malignant tumor.

 Nutritional supplements are often given in order to treat the symptoms of cachexia. This, however, often enhances the progress of the malignant tumor, and may shorten the survival time of the patient.

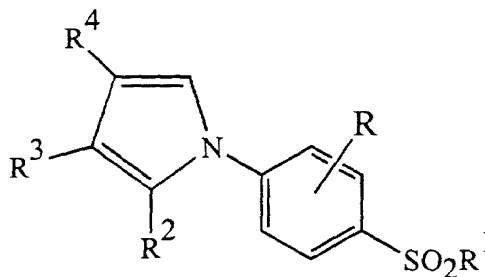
25 At present, no satisfactory treatment for cachexia has been established, and there is an increasing need for agents that alleviate the symptoms of cachexia.

 The compounds of formula (I) or (II), shown below, which, with certain other

compounds, are the active ingredients of the compositions of the present invention, are known to inhibit selectively cyclooxygenase-2 (COX-2). They are also known to inhibit the production of inflammatory cytokines (particularly IL-1 and TNF- α), to inhibit the production of leukotrienes (particularly LTB₄), to inhibit bone resorption, and to have analgesic, anti-inflammatory and anti-pyretic effects (European Patent Publication No. 799 823A).



(I)



(II)

It has not previously been known that these compounds can be used for the treatment or prevention of cachexia.

Also, although it is known that certain other active ingredients employed in the present invention, namely the compounds of formula (III), (IV), (V), (VI), (VII), (VIII), (IX), (X), (XI), (XII), (XIII) and (XIV) have selective inhibitory activity against cyclooxygenase-2, an inhibitory effect on the production of inflammatory cytokines (particularly IL-1 and TNF- α), an inhibitory action on the production of leukotrienes (particularly LTB₄), an inhibitory action on bone resorption, an analgesic action, an anti-inflammatory action and an antipyretic action [International publication number WO95/00501, J. Med. Chem., 40, 1347 (1997), International publication number WO94/13635, Pharmacology, 55, 44 (1997), Prostaglandins, 47, 55 (1994), Japanese publication number Hei 9-52882, Jpn. J. Pharmacol., 67,305 (1995), Inflamm. Res., 47, Suppl. 3, S257 (1997), J. Med. Chem., 38, 4570 (1995), US Patent No. 5 474 995, European Patent No. 863 134 and International Patent Publication No. WO 98/06708], it has not previously been disclosed that these compounds have an effect against cachexia.

It is known from epidemiological studies that the taking of conventional NSAIDS (non-steroidal anti-inflammatory drugs, which are COX-1 and COX-2 inhibitors), the most common of which is aspirin, and the incidence of colon cancer have an inverse correlation. In addition, there have been many reports that NSAIDS, such as aspirin and sulindac, have shown inhibitory activity against tumor metastasis and carcinogenesis in preclinical studies. Some NSAIDS have been used in clinical studies for the prevention of colon carcinogenesis.

However, since conventional NSAIDS are not selective for COX-1 or COX-2, the occurrence of adverse effects is unavoidable.

It would, therefore, be desirable to discover a selective cyclooxygenase-2 inhibitor (selective COX-2 inhibitor) for use as an anti-tumor agent that has a low level of adverse effects.

Among the known selective COX-2 inhibitors, it is known that MF-tricyclic [Oshima, M. *et al.* "Suppression of Intestinal Polyposis in APC Δ^{716} Knockout Mice by Inhibition of Cyclooxygenase 2 (COX-2)", *Cell*, 87, 803-809 (1996)] and celecoxib (Reddy, R.S. *et al.* "Evaluation of Cyclooxygenase-2 Inhibitor for Potential Chemopreventive Properties in Colon Carcinogenesis", *Cancer Res.*, 56, 4566-4569 (1996)] inhibit the occurrence of experimental colonic polyposis, and that SC-58125 exhibits growth inhibitory effects against certain types of human colon cancer cell lines (Sheng, H. *et al.* "Inhibition of Human Colon Cancer Cell Growth by Selective Inhibition of Cyclooxygenase-2", *J. Clin. Invest.*, 99, 2254-2259 (1997)].

However, in the case of the former, the experimental system used is not a model for an established colon cancer, and the compounds are only able to prevent the occurrence of polyposis in the preliminary stage of colon cancer.

On the other hand, with respect to the latter, the only colon cancer cell line in which growth inhibitory effects against human colon cancer cell lines have been observed is a cell line that expresses cyclooxygenase-2 (human colon cancer cell line HCA-7), and it has been disclosed that colon cancer cell lines that do not exhibit tumor growth inhibitory activity (HCT-116) *in vitro* do not exhibit tumor growth

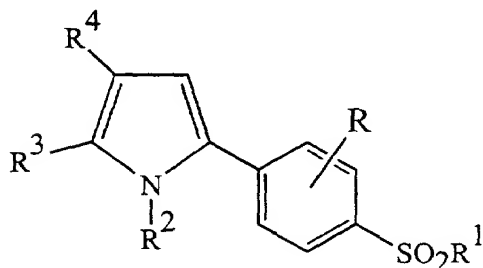
inhibitory effects *in vivo*. Thus, whether or not COX-2 inhibitor-induced tumor growth inhibitory effects on colon cancer are expressed *in vivo* depends on the sensitivity of the colon cancer cell lines used against COX-2 inhibitor-induced cell growth inhibitory activity *in vitro*. It is thus unlikely that the tumor growth
5 inhibitory effects of COX-2 inhibitors *in vivo* would be observed against various other cancers, especially those cancers, including colon cancers, that are resistant to COX-2 inhibitor-induced inhibition of cell growth *in vitro* and that do not express cyclooxygenase-2.

Moreover, there has been no previous disclosure of the use of a combination of
10 a selective cyclooxygenase-2 inhibitor and a 5-fluorouracil derivative for the prevention or inhibition of tumor growth.

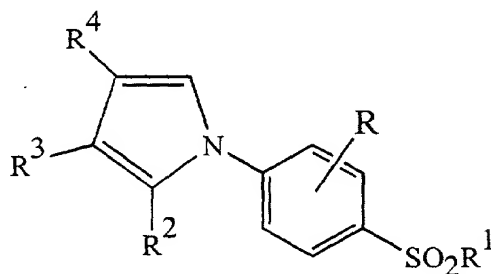
We have now found that certain 1,2-diphenylpyrrole derivatives and closely related compounds have excellent activity for the prevention or inhibition of cachexia, and that these 1,2-diphenylpyrrole derivatives are effective for the
15 treatment or prevention of tumor-related disorders, alone or in combination with a 5-fluorouracil derivative.

Brief Summary of the Invention

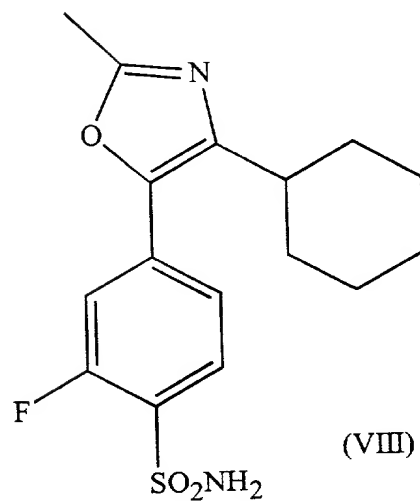
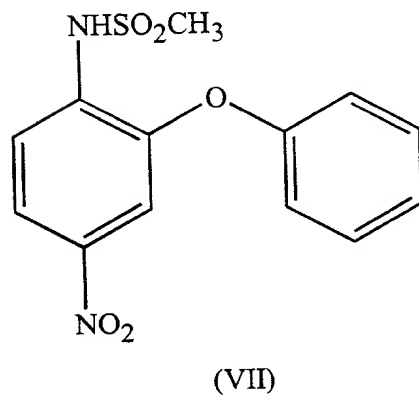
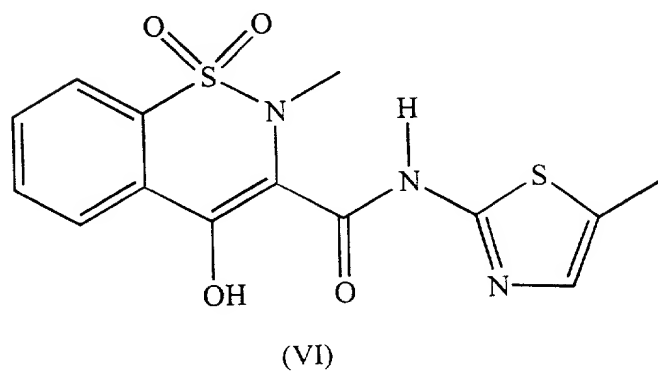
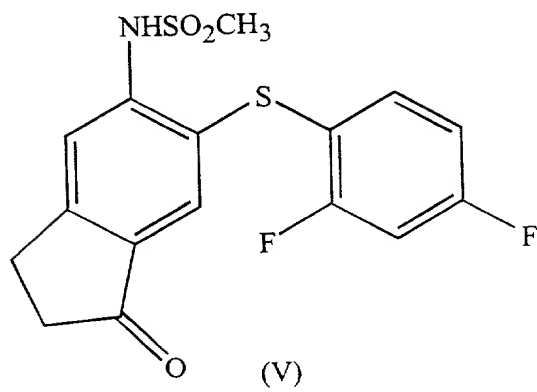
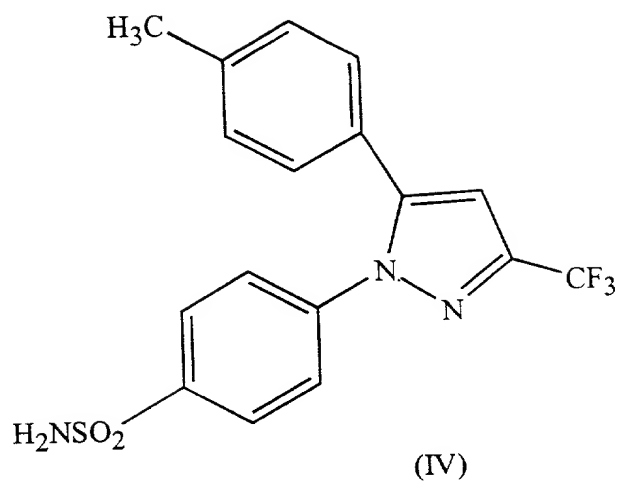
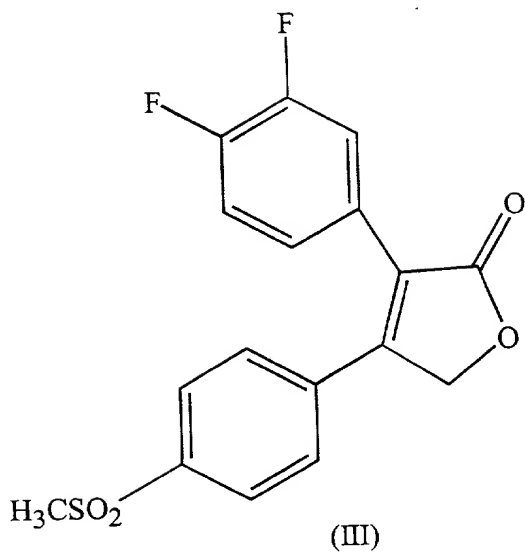
Thus, in a first embodiment, the present invention provides a method for the treatment or prevention of cachexia in a mammal, which may be human, in need of
20 such treatment or prevention, which method comprises administering to said mammal an effective amount of an active compound selected from the group consisting of compounds of formula (I), (II), (III), (IV), (V), (VI), (VII), (VIII), (IX), (X), (XI), (XII), (XIII) and (XIV):

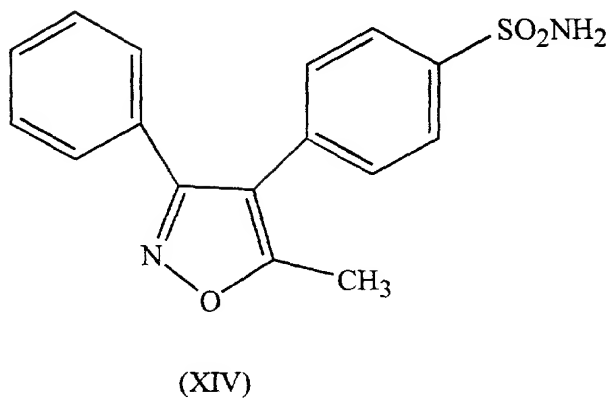
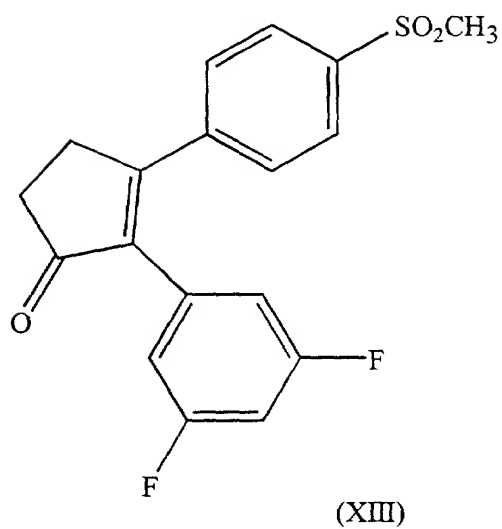
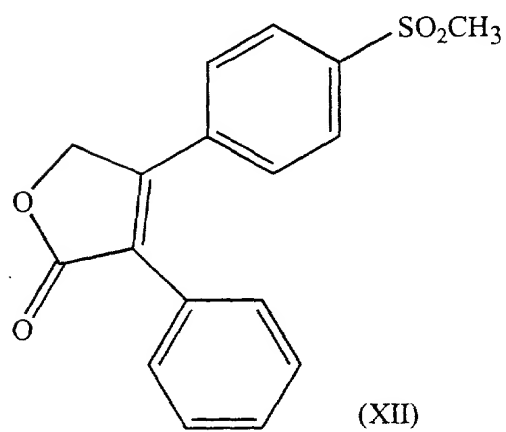
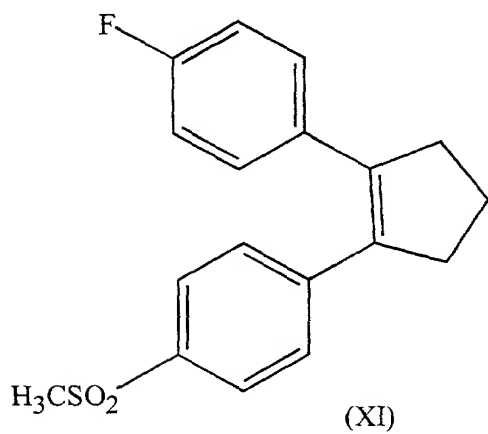
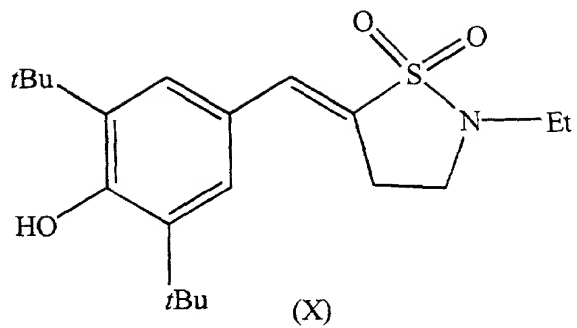
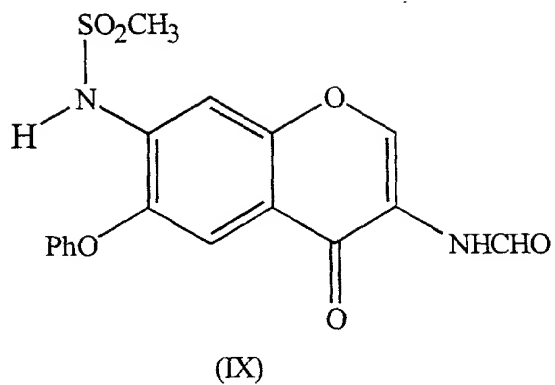


(I)



(II)





5

wherein

R represents a hydrogen atom, a halogen atom or a lower alkyl group;

R¹ represents a lower alkyl group, an amino group or a group of formula -
NHR^a (wherein R^a represents a group which may be eliminated *in vivo*);

5 R² represents a phenyl group or a phenyl group having at least one substituent
selected from the group consisting of substituents α and substituents β , defined
below;

R³ represents a hydrogen atom, a halogen atom, a lower alkyl group or a lower
alkyl group having at least one substituent selected from the group consisting
of substituents α ;

10 R⁴ represents a hydrogen atom, a lower alkyl group, a lower alkyl group
having at least one substituent selected from the group consisting of
substituents α , a cycloalkyl group, an aryl group as defined below, or an
aralkyl group as defined below;

15 said aryl group is a carbocyclic aromatic hydrocarbon group having from 6 to
14 carbon atoms in one or more aromatic rings or such a group which is fused
to a cycloalkyl group having from 3 to 10 carbon atoms, and the group is
unsubstituted or it is substituted by at least one substituent selected from the
group consisting of substituents α and substituents β ;

said aralkyl group is a lower alkyl group which is substituted by one or more of
the aryl groups defined above;

20 *t*Bu represents a t-butyl group;

Et represents an ethyl group; and

Ph represents a phenyl group;

said substituents α are selected from the group consisting of hydroxy groups,
halogen atoms, lower alkoxy groups and lower alkylthio groups; and

25 said substituents β are selected from the group consisting of lower alkyl
groups, alkanoyloxy groups, mercapto groups, alkanoylthio groups, lower

alkylsulfinyl groups, lower alkyl groups having at least one substituent selected from the group consisting of substituents α , cycloalkyloxy groups, lower haloalkoxy groups and lower alkylendioxy groups;

and pharmaceutically acceptable salts thereof.

- 5 The invention further provides a method for the treatment or prevention of tumor-related disorders in a mammal, which may be human, in need of such treatment or prevention, which method comprises administering to said mammal an effective amount of an active compound selected from the group consisting of compounds of formula (I) and (II), defined above.

10 **Detailed Description of Invention**

Preferred classes of compounds of the present invention are those compounds of formula (I) and (II) in which:

- (1) R represents a hydrogen atom, a fluorine atom, a chlorine atom or a methyl group, more preferably the hydrogen atom,
- 15 (2) R^1 represents a methyl group, an amino group or an acetylamino group, more preferably the amino group or acetylamino group,
- (3) R^2 represents a phenyl group or a phenyl group having at least one substituent selected from the group consisting of substituents α^1 and substituents β^1 , more preferably a phenyl group or phenyl group having at least one substituent selected
- 20 from the group consisting of substituents α^1 and substituents β^2 , still more preferably a phenyl group in which the number of substituents is from 1 to 3,
- (4) R^3 represents a hydrogen atom, a halogen atom, a lower alkyl group or a lower alkyl group having at least one substituent selected from the group consisting of substituents α^1 , more preferably a hydrogen atom, a halogen atom, a lower alkyl
- 25 group or a lower alkyl group substituted with a halogen atom,
- (5) R^4 represents a hydrogen atom, a lower alkyl group, a lower alkyl group having

at least one substituent selected from the group consisting of substituents α , a cycloalkyl group, an aryl group, an aryl group having at least one substituent selected from the group consisting of substituents α^1 and substituents β^3 , an aralkyl group or an aralkyl group having at least one substituent selected from the group consisting of substituents α^1 and substituents β^3 , more preferably a hydrogen atom, a lower alkyl group, a lower alkyl group having at least one substituent selected from the group consisting of substituents α^2 , a cycloalkyl group, an aryl group, an aryl group having at least one substituent selected from the group consisting of substituents α^2 and substituents β^4 , an aralkyl group or an aralkyl group having at least one substituent selected from the group consisting of substituents α^2 and substituents β^4 .

Said substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups.

Said substituents α^2 are selected from the group consisting of hydroxy groups, halogen atoms and lower alkoxy groups.

Said substituents β^1 are selected from the group consisting of lower alkyl groups, mercapto groups, alkanoylthio groups, lower alkyl groups having at least one substituent selected from the group consisting of substituents α^1 , lower haloalkoxy groups and lower alkylenedioxy groups.

Said substituents β^2 are selected from the group consisting of lower alkyl groups, mercapto groups, alkanoylthio groups, lower alkyl groups substituted with a halogen atom, lower haloalkoxy groups and lower alkylenedioxy groups.

Said substituents β^3 are selected from the group consisting of lower alkyl groups, lower alkyl groups having at least one substituent selected from the group consisting of substituents α and cycloalkyloxy groups.

Said substituents β^4 are selected from the group consisting of lower alkyl groups, lower alkyl groups substituted with a halogen atom and cycloalkyloxy

groups.

In the compounds of formula (I) and (II), where R, R³, substituents α , substituents α^1 or substituents α^2 represents a halogen atom, or where substituents β^2 or substituents β^4 represents a lower alkyl group substituted with halogen atom, the halogen atom is preferably a fluorine atom, a chlorine atom, a bromine atom or an iodine atom, preferably a fluorine atom, a chlorine atom or a bromine atom.

Where R, R¹, R³, R⁴, substituent β , substituent β^1 , substituent β^2 , substituent β^3 or substituent β^4 represents a lower alkyl group, or R³, R⁴, substituent β , substituent β^1 or substituent β^3 represents a lower alkyl group having at least one substituent selected from the group consisting of substituents α , or substituent β^2 or substituent β^4 represents a lower alkyl group substituted with a halogen atom, the alkyl group or alkyl part of the substituted group may be a straight or branched chain alkyl group having from 1 to 6 carbon atoms, and examples include the methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, t-butyl, pentyl, isopentyl, 2-methylbutyl, neopentyl, 1-ethylpropyl, hexyl, isohexyl, 4-methylpentyl, 3-methylpentyl, 2-methylpentyl, 1-methylpentyl, 3,3-dimethylbutyl, 2,2-dimethylbutyl, 1,1-dimethylbutyl, 1,2-dimethylbutyl, 1,3-dimethylbutyl, 2,3-dimethylbutyl and 2-ethylbutyl groups. Of these, we prefer the straight or branched chain alkyl groups having from 1 to 4 carbon atoms, more preferably the methyl and ethyl groups. In R, R¹ and R⁴, the lower alkyl group is particularly preferably the methyl group.

Where substituent β represents an alkanoyloxy group, or substituent β , substituent β^1 or substituent β^2 represents an alkanoylthio group, the alkanoyl part of these groups may be, for example, a straight or branched chain alkanoyl group having from 1 to 25 carbon atoms, such as the formyl, acetyl, propionyl, butyryl, isobutyryl, valeryl, isovaleryl, pivaloyl, hexanoyl, heptanoyl, octanoyl, nonanoyl, decanoyl, undecanoyl, lauroyl, tridecanoyl, myristoyl, palmitoyl, stearoyl, icosanoyl, docosanoyl and pentacosanoyl groups. Of these, we prefer those alkanoyl groups having from 1 to 12 carbon atoms, more preferably those alkanoyl groups having from 1 to 6 carbon atoms, still more preferably those alkanoyl groups having from 1

to 4 carbon atoms, and most preferably the acetyl and propionyl groups.

Where R^4 represents a cycloalkyl group, this is preferably a cycloalkyl group having from 3 to 8 carbon atoms, such as the cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl and cyclooctyl groups. Of these, we prefer those cycloalkyl
5 group having from 3 to 7 carbon atoms, more preferably those cycloalkyl groups having from 3 to 6 carbon atoms, and most preferably the cyclopropyl group.

Where R^4 represents an aryl group, this aryl group may be a carbocyclic aromatic hydrocarbon group having from 6 to 14 carbon atoms and may be unsubstituted or it may be substituted by at least one substituent selected from the
10 group consisting of substituents α and substituents β . The group may contain a single aromatic ring or it may contain two or more fused rings. Examples of such groups include the phenyl, indenyl, naphthyl, phenanthrenyl and anthracenyl groups. Of these, we prefer the phenyl and naphthyl groups, more preferably the phenyl group. The above-mentioned aryl group may be condensed with a cycloalkyl group
15 having from 3 to 10 carbon atoms and examples of such condensed groups include, for example, the 2-indanyl group.

Where R^4 represents an aralkyl group, this is an alkyl group, which may be any of the alkyl groups defined and exemplified above in relation to R etc., and which is substituted by from 1 to 3 aryl groups, as defined and exemplified above. Such a
20 group may be unsubstituted or it may be substituted by at least one of substituents α or β . Examples of such groups include the benzyl, phenethyl, 3-phenylpropyl, 4-phenylbutyl, 1-naphthylmethyl, 2-naphthylmethyl, diphenylmethyl, triphenylmethyl, 1-naphthylmethyl and 9-anthrylmethyl groups. Of these, we prefer an alkyl group having from 1 to 4 carbon atoms which is substituted with one aryl group
25 having from 6 to 10 carbon atoms.

Where substituent α , substituent α^1 or substituent α^2 represents a lower alkoxy group, this may be, for example, a straight or branched chain alkoxy group having from 1 to 6 carbon atoms, such as the methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutoxy, sec-butoxy, t-butoxy, pentyloxy, isopentyloxy, 2-methylbutoxy,

neopentyloxy, 1-ethylpropoxy, hexyloxy, isohexyloxy, 4-methylpentyloxy, 3-methylpentyloxy, 2-methylpentyloxy, 1-methylpentyloxy, 3,3-dimethylbutoxy, 2,2-dimethylbutoxy, 1,1-dimethylbutoxy, 1,2-dimethylbutoxy, 1,3-dimethylbutoxy, 2,3-dimethylbutoxy or 2-ethylbutoxy groups. Of these, we prefer the straight or
5 branched chain alkoxy groups having from 1 to 4 carbon atoms, more preferably the methoxy and ethoxy groups.

Where substituent α or substituent α^1 represents a lower alkylthio group, this may be a straight or branched chain alkylthio group having from 1 to 6 carbon atoms, and examples include the methylthio, ethylthio, propylthio, isopropylthio, butylthio,
10 isobutylthio, sec-butylthio, t-butylthio, pentylthio, isopentylthio, 2-methylbutylthio, neopentylthio, 1-ethylpropylthio, hexylthio, isohexylthio, 4-methylpentylthio, 3-methylpentylthio, 2-methylpentylthio, 1-methylpentylthio, 3,3-dimethylbutylthio, 2,2-dimethylbutylthio, 1,1-dimethylbutylthio, 1,2-dimethylbutylthio, 1,3-dimethylbutylthio, 2,3-dimethylbutylthio and 2-ethylbutylthio groups. Of these, we
15 prefer the straight or branched chain alkylthio groups having from 1 to 4 carbon atoms, more preferably the methylthio and ethylthio groups.

Where substituent β represents a lower alkylsulfinyl group, this may be a straight or branched chain alkylsulfinyl group having from 1 to 6 carbon atoms, such as the methylsulfinyl, ethylsulfinyl, propylsulfinyl, isopropylsulfinyl, butylsulfinyl,
20 isobutylsulfinyl, sec-butylsulfinyl, t-butylsulfinyl, pentylsulfinyl, isopentylsulfinyl, 2-methylbutylsulfinyl, neopentylsulfinyl, 1-ethylpropylsulfinyl, hexylsulfinyl, isohexylsulfinyl, 4-methylpentylsulfinyl, 3-methylpentylsulfinyl, 2-methylpentylsulfinyl, 1-methylpentylsulfinyl, 3,3-dimethylbutylsulfinyl, 2,2-dimethylbutylsulfinyl, 1,1-dimethylbutylsulfinyl, 1,2-dimethylbutylsulfinyl, 1,3-dimethylbutylsulfinyl, 2,3-dimethylbutylsulfinyl and 2-ethylbutylsulfinyl groups. Of
25 these, we prefer the straight or branched chain alkylsulfinyl groups having from 1 to 4 carbon atoms.

Where substituent β , substituent β^3 or substituent β^4 represents a cycloalkyloxy group, this may be, for example, a cycloalkyloxy group having from 3
30 to 8 carbon atoms, such as the cyclopropyloxy, cyclobutyloxy, cyclopentyloxy,

cyclohexyloxy, cycloheptyloxy and cyclooctyloxy groups. Of these, we prefer the cycloalkyloxy groups having from 3 to 7 carbon atoms, more preferably the cycloalkyloxy groups having 5 or 6 carbon atoms, most preferably the cyclopentyloxy group.

5 Where substituent β , substituent β^1 or substituent β^2 represents a lower haloalkoxy group, this is an alkoxy group, which may be as defined and exemplified above in relation to substituent α etc., and which is substituted by at least one halogen atom, such as those defined and exemplified above. Examples of such groups include the fluoromethoxy, difluoromethoxy, trifluoromethoxy, 2-
10 fluoroethoxy, 2-chloroethoxy, 2-bromoethoxy, 2,2-difluoroethoxy, 2,2,2-trifluoroethoxy, 2,2,2-trichloroethoxy, 3-fluoropropoxy, 4-fluorobutoxy, chloromethoxy, trichloromethoxy, iodomethoxy and bromomethoxy groups. Of these, we prefer those lower haloalkoxy groups having from 1 to 4 carbon atoms, more preferably the fluoromethoxy, difluoromethoxy, trifluoromethoxy, 2-
15 fluoroethoxy, 2-chloroethoxy, 2-bromoethoxy, 3-fluoropropoxy, 4-fluorobutoxy, chloromethoxy, trichloromethoxy and bromomethoxy groups, and most preferably the fluoromethoxy, difluoromethoxy and trifluoromethoxy groups.

 Where substituent β , substituent β^1 or substituent β^2 represents a lower alkylendioxy group, this may be, for example, a straight or branched chain
20 alkylendioxy group having from 1 to 6 carbon atoms, such as the methylenedioxy, ethylenedioxy, trimethylenedioxy, tetramethylenedioxy, pentamethylenedioxy, hexamethylenedioxy and propylenedioxy groups. Of these, we prefer those alkylendioxy groups having from 1 to 4 carbon atoms, more preferably the methylenedioxy and ethylenedioxy groups.

25 Where substituent β^2 or substituent β^4 represents a lower alkyl group substituted with a halogen atom, this may be any of the alkyl groups defined and exemplified above in relation to R etc., which is substituted by at least one halogen atom, as also defined and exemplified above. Examples of such groups include the fluoromethyl, difluoromethyl, trifluoromethyl, 2-fluoroethyl, 2,2-difluoroethyl, 2,2,2-
30 trifluoroethyl, 2,2,2-trichloroethyl, 3-fluoropropyl, 4-fluorobutyl, chloromethyl,

trichloromethyl, 2-chloroethyl, 3-chloropropyl, bromomethyl, 2-bromoethyl, iodomethyl, 2-iodoethyl, chlorodifluoromethyl and bromodifluoromethyl groups. Of these, we prefer those haloalkyl groups having from 1 to 4 carbon atoms, more preferably the fluoromethyl, difluoromethyl, trifluoromethyl, 2-fluoroethyl, 3-fluoropropyl, 4-fluorobutyl, chloromethyl, trichloromethyl and bromomethyl groups, and most preferably the fluoromethyl, difluoromethyl and trifluoromethyl groups.

Where R^a represents a group to be eliminated *in vivo* is a group which can be eliminated in the human body under physiological conditions such as hydrolysis, that is a group producing a free amino group ($-NH_2$) from a group of formula $-NHR^a$ (wherein R^a is as defined above). It is easy to determine whether or not the group can be eliminated *in vivo* by the following test: a compound to be tested is administered orally or intravenously to an experimental animal, such as a rat or mouse, and the body fluid is tested for the presence or absence of the corresponding compound having a free amino group or a pharmaceutically acceptable salt thereof.

Such groups include, for example:

the alkanoyl groups defined and exemplified above in relation to substituent β etc.;

a lower alkoxycarbonyl group, in which the alkoxy group is as defined and exemplified above in relation to substituent α etc., such as the methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, isopropoxycarbonyl, butoxycarbonyl, isobutoxycarbonyl, t-butoxycarbonyl, pentyloxycarbonyl, hexyloxycarbonyl and cyclohexyloxycarbonyl groups;

an aralkyloxycarbonyl group in which the aryl is as defined above and is unsubstituted or is substituted by one or two lower alkoxy or nitro groups, such as the benzyloxycarbonyl, 4-methoxybenzyloxycarbonyl, 3,4-dimethoxybenzyloxycarbonyl, 2-nitrobenzyloxycarbonyl and 4-nitrobenzyloxycarbonyl groups;

an alkanoyloxymethyl group in which the alkanoyl group is as defined and exemplified above in relation to substituent β etc., such as the formyloxy-

methyl, acetoxymethyl, propionyloxymethyl, butyryloxymethyl, pivaloyloxymethyl, valeryloxymethyl, isovaleryloxymethyl and hexanoyloxymethyl groups;

5 a lower alkoxy-carbonyloxymethyl group in which the alkoxy group is as defined and exemplified above in relation to substituent α etc., such as the methoxycarbonyloxymethyl, ethoxycarbonyloxymethyl, propoxycarbonyloxymethyl, isopropoxycarbonyloxymethyl, butoxycarbonyloxymethyl, isobutoxycarbonyloxymethyl and pentyloxycarbonyloxymethyl groups; and

10 a (2-oxo-1,3-dioxolen-4-yl)methyl group in which the 5-position of the dioxolene ring may be substituted with a lower alkyl group or an aryl group, as defined and exemplified above in relation to R and R^4 , respectively, such as the (5-phenyl-2-oxo-1,3-dioxolen-4-yl)methyl, [5-(4-methylphenyl)-2-oxo-1,3-dioxolen-4-yl]methyl, [5-(4-methoxyphenyl)-2-oxo-1,3-dioxolen-4-yl]methyl, [5-(4-fluorophenyl)-2-oxo-1,3-dioxolen-4-yl]methyl, [5-(4-chlorophenyl)-2-oxo-1,3-dioxolen-4-yl]methyl, (2-oxo-1,3-dioxolen-4-yl)methyl, (5-methyl-2-oxo-1,3-dioxolen-4-yl)methyl, (5-ethyl-2-oxo-1,3-dioxolen-4-yl)methyl, (5-propyl-2-oxo-1,3-dioxolen-4-yl)methyl, (5-isopropyl-2-oxo-1,3-dioxolen-4-yl)methyl and (5-butyl-2-oxo-1,3-dioxolen-4-yl)methyl groups.

Of these, we prefer the alkanoyl groups having from 1 to 12 carbon atoms, the 20 alkoxy-carbonyl groups having from 2 to 5 carbon atoms, the aralkyloxycarbonyl groups having 7 to 8 carbon atoms, the alkanoyloxymethyl groups having from 3 to 6 carbon atoms, the alkoxy-carbonyloxymethyl groups having from 3 to 6 carbon atoms and the 5-substituted (2-oxo-1,3-dioxolen-4-yl)methyl group, more preferably the acetyl, propionyl, butyryl, isobutyryl, valeryl, isovaleryl, pivaloyl, methoxycarbonyl, 25 ethoxycarbonyl, benzyloxycarbonyl, acetoxymethyl, propionyloxymethyl, methoxycarbonyloxymethyl, ethoxycarbonyloxymethyl, (5-methyl-2-oxo-1,3-dioxolen-4-yl)methyl and (5-phenyl-2-oxo-1,3-dioxolen-4-yl)methyl groups, and most preferably the acetyl group.

Specific examples of R^1 preferably include the methyl, ethyl, amino, 30 acetylamino, propionylamino, butyrylamino, isobutyrylamino, valerylamino,

isovalerylamino, pivaloylamino, methoxycarbonylamino, ethoxycarbonylamino, benzyloxycarbonylamino, acetoxymethylamino, propionyloxymethylamino, methoxycarbonyloxymethylamino, ethoxycarbonyloxymethylamino, (5-methyl-2-oxo-1,3-dioxolen-4-yl)methylamino and (5-phenyl-2-oxo-1,3-dioxolen-4-yl)methyl-
5 amino groups, more preferably the methyl, amino and acetylamino groups, and most preferably the amino and acetylamino groups.

Specific examples of R^2 preferably include:

the unsubstituted phenyl group;

10 phenyl groups having from 1 to 3 substituents selected from mercapto groups, $C_1 - C_4$ alkanoylthio groups, halogen atoms, $C_1 - C_4$ alkyl groups, $C_1 - C_4$ alkoxy groups, $C_1 - C_4$ alkylthio groups and $C_1 - C_4$ alkylsulfinyl groups, such as the 4-mercaptophenyl, 4-acetylthiophenyl, 4-propionylthiophenyl, 4-fluorophenyl, 4-chlorophenyl, 4-bromophenyl, p-tolyl, 4-ethylphenyl, 4-methoxyphenyl, 4-ethoxyphenyl, 4-methylthiophenyl, 4-ethylthiophenyl, 4-methylsulfinylphenyl, 4-ethylsulfinylphenyl, 3,4-difluorophenyl, 2,4-difluorophenyl, 3,4-dichlorophenyl, 2,4-dichlorophenyl, 3,4-dimethylphenyl, 3,4-dimethoxyphenyl, 3-chloro-4-fluorophenyl, 3-chloro-4-methoxyphenyl, 3-fluoro-4-methoxyphenyl, 3-methyl-4-methoxyphenyl, 3,5-dichloro-4-methoxyphenyl and 4-methoxy-3,5-dimethylphenyl groups;

20 trifluoromethyl-, difluoromethoxy- or trifluoromethoxy-substituted phenyl groups, such as the 4-trifluoromethylphenyl, 4-difluoromethoxyphenyl and 4-trifluoromethoxyphenyl groups;

methylenedioxy- or ethylenedioxy-substituted phenyl group such as the 3,4-methylenedioxyphenyl and 3,4-ethylenedioxyphenyl groups.

25 In the case where R^2 is a substituted phenyl group, the number of substituents is preferably from 1 to 3, more preferably 1 or 2.

Specific examples of R^3 preferably include hydrogen atoms; halogen atoms,

such as the fluorine, chlorine, bromine and iodine atoms; C₁ - C₄ alkyl groups, such as the methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl and t-butyl groups; and C₁ - C₄ haloalkyl groups, such as the fluoromethyl, chloromethyl, bromomethyl, iodomethyl, difluoromethyl, trifluoromethyl, 2-fluoroethyl, 3-fluoropropyl, 4-fluorobutyl, 2-chloroethyl and 3-chloropropyl groups, more preferably hydrogen atoms; halogen atoms, such as the fluorine, chlorine, bromine and iodine atoms; and the methyl, ethyl, fluoromethyl, difluoromethyl, 2-fluoroethyl and 2-chloroethyl groups.

Specific examples of R⁴ preferably include hydrogen atoms; C₁ - C₆ alkyl groups, such as the methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, t-butyl, pentyl and hexyl groups; any of these alkyl groups having a substituent selected from hydroxy, halogen (such as fluorine, chlorine, bromine or iodine) and C₁ - C₄ alkoxy (such as methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutoxy, sec-butoxy and t-butoxy); C₃ - C₇ cycloalkyl groups, such as the cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl and cycloheptyl groups; C₆ - C₁₀ aryl groups, such as the phenyl and naphthyl groups, which may be unsubstituted or may have one or more of the following substituents γ; C₆ - C₁₀ aryl C₁ - C₄ alkyl groups, such as the benzyl, phenethyl, 3-phenylpropyl, 4-phenylbutyl, 1-naphthylmethyl and 2-naphthylmethyl groups, which may be unsubstituted or may have one or more of the following substituents γ in the aryl moiety;

substituents γ include: halogen atoms, such as the fluorine, chlorine, bromine and iodine atoms; C₁ - C₄ alkyl groups, such as the methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl and t-butyl groups; C₁ - C₄ haloalkyl groups, such as the fluoromethyl, difluoromethyl, trifluoromethyl, chloromethyl, trichloromethyl, chlorodifluoromethyl, 2-fluoroethyl, 2-chloroethyl, 2-bromoethyl, 2-iodoethyl, 3-fluoropropyl and 4-fluoropropyl groups; C₁ - C₄ alkoxy groups, such as the methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutoxy, sec-butoxy and t-butoxy groups; and C₃ - C₇ cycloalkyloxy groups, such as the cyclopropyloxy, cyclobutyloxy, cyclopentyloxy, cyclohexyloxy and cycloheptyloxy groups.

Preferred examples of R⁴ include: hydrogen atoms; C₁ - C₄ alkyl groups, such as the methyl, ethyl, isopropyl, butyl and isobutyl groups; C₁ - C₄ mono-, di- or trihaloalkyl groups, such as the fluoromethyl, difluoromethyl, chlorodifluoromethyl, bromodifluoromethyl, trifluoromethyl, 2-fluoroethyl and 2,2,2-trifluoroethyl groups; 5 hydroxymethyl groups; C₁ - C₄ alkoxymethyl groups, such as the methoxymethyl and ethoxymethyl groups; C₃ - C₆ cycloalkyl groups, such as the cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl groups; phenyl groups; mono- or difluorophenyl groups, such as the 4-fluorophenyl and 2,4-difluorophenyl groups; mono- or dimethoxyphenyl groups, such as the 4-methoxyphenyl and 3,4-dimethoxyphenyl 10 groups; tolyl groups, such as the *p*-tolyl and *o*-tolyl groups; cyclopentyloxy-(methoxy)phenyl groups, such as the 3-cyclopentyloxy-4-methoxyphenyl group; trifluoromethylphenyl groups, such as the 4-trifluoromethylphenyl group; benzyl groups; substituted benzyl groups, such as the 4-methoxybenzyl and 3-cyclopentyloxy-4-methoxybenzyl groups; phenethyl groups; naphthyl groups, such as the 15 1-naphthyl and 2-naphthyl groups; and naphthylmethyl groups, such as the 1-naphthylmethyl and 2-naphthylmethyl groups.

Certain of the compounds of the present invention, specifically the compounds of formula (I) and (II), possess an acidic group and can thus form salts with cations. The nature of the salt is not critical to the present invention, provided that it is 20 pharmaceutically acceptable, that is that the salt is neither less active (or unacceptably less active) nor more toxic (or unacceptably more toxic) than the free acid. Such salts include, for example: salts with alkali metals, such as sodium, potassium or lithium; salts with alkaline earth metals, such as calcium or magnesium; salts with other metals, such as aluminum, iron, zinc, copper, nickel or cobalt; other 25 inorganic salts, such as the ammonium salt; salts with organic amines, such as *t*-octylamine, dibenzylamine, morpholine, glucosamine, phenylglycine alkyl ester, ethylenediamine, *N*-methylglucamine, guanidine, diethylamine, triethylamine, dicyclohexylamine, *N,N'*-dibenzylethylenediamine, chlorprocaine, procaine, diethanolamine, *N*-benzyl-*N*-phenethylamine, piperazine, tetramethyl ammonium or 30 tris(hydroxymethyl)aminomethane.

Further, when the compounds of formula (I) and (II) and salts thereof are allowed to stand in the atmosphere, they may adsorb moisture to form hydrates. Such hydrates are also included in the present invention.

Further, the compounds of formula (I) and (II) and salts thereof sometimes
5 absorb certain kinds of solvents to afford solvates, and such solvates are also included in the present invention.

Certain of the compounds of formula (I) and (II) of the present invention may have asymmetric carbon atoms in their molecule, and stereoisomers in the R-configuration or the S-configuration would then exist. Each of these stereoisomers
10 and mixtures thereof in any desired proportion are all included in the present invention.

Specific examples of the compounds of formula (I) and (II) employed in the method and composition of the present invention include, for example, those shown in the following Table 1 [compounds of formula (I)] and Table 2 [compounds of
15 formula (II)].

In the Tables, the following abbreviations are used:

	Ac	acetyl;
	Bu	butyl;
	Byr	butyryl;
20	iByr	isobutyryl;
	Bz	benzyl;
	Et	ethyl;
	For	formyl;
	Me	methyl;
25	Ph	phenyl;
	Piv	pivaloyl;
	cPn	cyclopentyl;
	Pr	propyl;
	cPr	cyclopropyl;
30	iPr	isopropyl;

Pm	propionyl;
iVal	isovaleryl; and
Val	valeryl.

Table 1

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
1-1	H	Me	Ph	H	H
1-2	H	Me	Ph	H	Me
1-3	H	Me	4-F-Ph	H	H
1-4	H	Me	4-F-Ph	F	H
1-5	H	Me	4-F-Ph	Cl	H
1-6	H	Me	4-F-Ph	Br	H
1-7	H	Me	4-F-Ph	I	H
1-8	H	Me	4-F-Ph	Me	H
1-9	H	Me	4-F-Ph	Et	H
1-10	H	Me	4-F-Ph	Pr	H
1-11	H	Me	4-F-Ph	Bu	H
1-12	H	Me	4-F-Ph	CH ₂ F	H
1-13	H	Me	4-F-Ph	CHF ₂	H
1-14	H	Me	4-F-Ph	CF ₃	H
1-15	H	Me	4-F-Ph	H	Me
1-16	H	Me	4-F-Ph	F	Me
1-17	H	Me	4-F-Ph	Cl	Me
1-18	H	Me	4-F-Ph	Br	Me
1-19	H	Me	4-F-Ph	I	Me
1-20	H	Me	4-F-Ph	Me	Me

Table 1 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
1-21	H	Me	4-F-Ph	Et	Me
1-22	H	Me	4-F-Ph	Pr	Me
1-23	H	Me	4-F-Ph	H	Et
1-24	H	Me	4-F-Ph	H	Pr
1-25	H	Me	4-F-Ph	H	Bu
1-26	H	Me	4-F-Ph	H	cPr
1-27	H	Me	4-F-Ph	H	Ph
1-28	H	Me	4-F-Ph	H	CH ₂ Ph
1-29	H	Me	4-F-Ph	H	CHF ₂
1-30	H	Me	4-F-Ph	Me	CHF ₂
1-31	H	Me	4-F-Ph	H	CF ₃
1-32	H	Me	4-F-Ph	Me	CF ₃
1-33	H	Me	4-MeO-Ph	H	H
1-34	H	Me	4-MeO-Ph	H	Me
1-35	H	Me	4-Cl-Ph	H	H
1-36	H	Me	4-Cl-Ph	H	Me
1-37	H	Me	4-Me-Ph	H	H
1-38	H	Me	4-Me-Ph	H	Me
1-39	H	Me	3-Cl-4-F-Ph	H	H
1-40	H	Me	3-Cl-4-F-Ph	H	Me

Table 1 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
1-41	H	Me	3,4-methylenedioxy-Ph	H	H
1-42	H	Me	3,4-methylenedioxy-Ph	H	Me
1-43	H	Me	3-Cl-4-MeO-Ph	H	H
1-44	H	Me	3-Cl-4-MeO-Ph	H	Me
1-45	H	Me	4-CF ₃ -Ph	H	H
1-46	H	Me	4-CF ₃ O-Ph	H	H
1-47	H	Me	3-F-4-MeO-Ph	H	H
1-48	H	Me	3-F-4-MeO-Ph	H	Me
1-49	H	Me	3-Me-4-MeO-Ph	H	H
1-50	H	Me	3-Me-4-MeO-Ph	H	Me
1-51	H	Me	3,4-diF-Ph	H	H
1-52	H	Me	3,4-diF-Ph	H	Me
1-53	H	Me	2,4-diF-Ph	H	H
1-54	H	Me	2,4-diF-Ph	H	Me
1-55	H	Me	3,4-diMe-Ph	H	H
1-56	H	Me	3,4-diMe-Ph	H	Me
1-57	H	Me	3,4-diCl-Ph	H	H
1-58	H	Me	3,4-diCl-Ph	H	Me
1-59	H	Me	3,4-di(MeO)-Ph	H	H
1-60	H	Me	3,4-di(MeO)-Ph	H	Me

Table 1 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
1-61	H	Me	4-F-Ph	H	CH ₂ OH
1-62	H	Me	4-F-Ph	Me	CH ₂ OH
1-63	H	Me	4-F-Ph	H	CH ₂ OMe
1-64	H	Me	4-MeO-Ph	H	CH ₂ OH
1-65	H	Me	4-MeO-Ph	H	CH ₂ OMe
1-66	H	Me	4-Cl-Ph	H	CH ₂ OH
1-67	H	Me	4-Cl-Ph	H	CH ₂ OMe
1-68	H	Me	4-Me-Ph	H	CH ₂ OH
1-69	H	Me	4-Me-Ph	H	CH ₂ OMe
1-70	H	NH ₂	Ph	H	H
1-71	H	NH ₂	Ph	H	Me
1-72	H	NH ₂	Ph	Me	H
1-73	H	NH ₂	4-F-Ph	H	H
1-74	H	NH ₂	4-F-Ph	H	Me
1-75	H	NH ₂	4-F-Ph	Cl	Me
1-76	H	NH ₂	4-F-Ph	Me	H
1-77	H	NH ₂	4-F-Ph	H	Et
1-78	H	NH ₂	4-F-Ph	H	Pr

Table 1 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
1-79	H	NH ₂	4-F-Ph	H	Bu
1-80	H	NH ₂	4-F-Ph	H	cPr
1-81	H	NH ₂	4-F-Ph	H	Ph
1-82	H	NH ₂	4-F-Ph	H	CH ₂ Ph
1-83	H	NH ₂	4-F-Ph	H	CHF ₂
1-84	H	NH ₂	4-F-Ph	H	CF ₃
1-85	H	NH ₂	4-MeO-Ph	H	H
1-86	H	NH ₂	4-MeO-Ph	H	Me
1-87	H	NH ₂	4-MeO-Ph	H	Bu
1-88	H	NH ₂	4-MeO-Ph	Me	H
1-89	H	NH ₂	4-EtO-Ph	H	H
1-90	H	NH ₂	4-EtO-Ph	H	Me
1-91	H	NH ₂	4-EtO-Ph	Me	H
1-92	H	NH ₂	4-PrO-Ph	H	Me
1-93	H	NH ₂	4-MeS-Ph	H	H
1-94	H	NH ₂	4-MeS-Ph	H	Me
1-95	H	NH ₂	4-MeS-Ph	Me	H
1-96	H	NH ₂	4-Cl-Ph	H	H

Table 1 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
1-97	H	NH ₂	4-Cl-Ph	H	Me
1-98	H	NH ₂	4-Cl-Ph	Me	H
1-99	H	NH ₂	4-Me-Ph	H	H
1-100	H	NH ₂	4-Me-Ph	H	Me
1-101	H	NH ₂	4-Me-Ph	Me	H
1-102	H	NH ₂	3-Cl-4-F-Ph	H	H
1-103	H	NH ₂	3-Cl-4-F-Ph	H	Me
1-104	H	NH ₂	3-Cl-4-F-Ph	Me	H
1-105	H	NH ₂	3,4-methylenedioxy-Ph	H	H
1-106	H	NH ₂	3,4-methylenedioxy-Ph	H	Me
1-107	H	NH ₂	3-Cl-4-MeO-Ph	H	H
1-108	H	NH ₂	3-Cl-4-MeO-Ph	H	Me
1-109	H	NH ₂	3-Cl-4-MeO-Ph	Me	H
1-110	H	NH ₂	4-CF ₃ -Ph	H	H
1-111	H	NH ₂	4-CF ₃ O-Ph	H	H
1-112	H	NH ₂	3-F-4-MeO-Ph	H	H
1-113	H	NH ₂	3-F-4-MeO-Ph	H	Me
1-114	H	NH ₂	3-F-4-MeO-Ph	Me	H

Table 1 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
1-115	H	NH ₂	3-Me-4-MeO-Ph	H	H
1-116	H	NH ₂	3-Me-4-MeO-Ph	H	Me
1-117	H	NH ₂	3-Me-4-MeO-Ph	Me	H
1-118	H	NH ₂	3,4-diF-Ph	H	H
1-119	H	NH ₂	3,4-diF-Ph	H	Me
1-120	H	NH ₂	3,4-diF-Ph	Me	H
1-121	H	NH ₂	2,4-diF-Ph	H	H
1-122	H	NH ₂	2,4-diF-Ph	H	Me
1-123	H	NH ₂	2,4-diF-Ph	Me	H
1-124	H	NH ₂	3,4-diMe-Ph	H	H
1-125	H	NH ₂	3,4-diMe-Ph	H	Me
1-126	H	NH ₂	3,4-diMe-Ph	Me	H
1-127	H	NH ₂	2,4-diCl-Ph	H	H
1-128	H	NH ₂	2,4-diCl-Ph	H	Me
1-129	H	NH ₂	2,4-diCl-Ph	Me	H
1-130	H	NH ₂	3,4-diCl-Ph	H	H
1-131	H	NH ₂	3,4-diCl-Ph	H	Me
1-132	H	NH ₂	3,4-diCl-Ph	Me	H

Table 1 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
1-133	H	NH ₂	3,4-di(MeO)-Ph	H	H
1-134	H	NH ₂	3,4-di(MeO)-Ph	H	Me
1-135	H	NH ₂	4-F-Ph	H	CH ₂ OH
1-136	H	NH ₂	4-F-Ph	H	CH ₂ OMe
1-137	H	NH ₂	4-MeO-Ph	H	CH ₂ OH
1-138	H	NH ₂	4-MeO-Ph	H	CH ₂ OMe
1-139	H	NH ₂	4-Cl-Ph	H	CH ₂ OH
1-140	H	NH ₂	4-Cl-Ph	H	CH ₂ OMe
1-141	H	NH ₂	4-Me-Ph	H	CH ₂ OH
1-142	H	NH ₂	4-Me-Ph	H	CH ₂ OMe
1-143	H	NH ₂	4-Et-Ph	H	H
1-144	H	NH ₂	4-Et-Ph	H	Me
1-145	H	NH ₂	4-Et-Ph	Me	H
1-146	H	NH ₂	2,4,6-triMe-Ph	H	Me
1-147	H	NH ₂	4-MeO-Ph	Cl	H
1-148	H	NH ₂	4-MeO-Ph	Br	H
1-149	H	NH ₂	4-MeO-Ph	Cl	Me
1-150	H	NH ₂	2-F-4-Cl-Ph	H	Me

Table 1 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
1-151	H	NH ₂	4-EtO-Ph	Cl	H
1-152	H	NH ₂	4-MeS-Ph	Cl	H
1-153	H	NH ₂	4-MeSO-Ph	H	Me
1-154	H	NH ₂	4-EtS-Ph	H	Me
1-155	H	NH ₂	2,4-diCl-Ph	Cl	H
1-156	H	NH ₂	4-SH-Ph	H	Me
1-157	H	NH ₂	4-AcS-Ph	H	Me
1-158	3-F	NH ₂	4-MeO-Ph	H	Me
1-159	3-F	NH ₂	4-EtO-Ph	H	Me
1-160	3-F	NH ₂	3,4-diMe-Ph	H	Me
1-161	3-F	NH ₂	4-Cl-Ph	H	Me
1-162	3-F	NH ₂	4-F-Ph	H	Me
1-163	3-F	NH ₂	4-SH-Ph	H	Me
1-164	3-F	NH ₂	4-MeS-Ph	H	Me
1-165	3-F	NH ₂	4-EtS-Ph	H	Me
1-166	3-F	NH ₂	4-AcS-Ph	H	Me
1-167	3-Me	NH ₂	4-MeO-Ph	H	Me
1-168	3-Me	NH ₂	4-EtO-Ph	H	Me

Table 1 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
1-169	3-Me	NH ₂	3,4-diMe-Ph	H	Me
1-170	3-Me	NH ₂	4-Cl-Ph	H	Me
1-171	3-Me	NH ₂	4-F-Ph	H	Me
1-172	3-Me	NH ₂	4-MeS-Ph	H	Me
1-173	H	NHFor	4-MeS-Ph	H	Me
1-174	H	NHAc	4-MeS-Ph	H	Me
1-175	H	NHPm	4-MeS-Ph	H	Me
1-176	H	NHByr	4-MeS-Ph	H	Me
1-177	H	NHiByr	4-MeS-Ph	H	Me
1-178	H	NHVal	4-MeS-Ph	H	Me
1-179	H	NHiVal	4-MeS-Ph	H	Me
1-180	H	NHPiv	4-MeS-Ph	H	Me
1-181	H	NH(MeOCO)	4-MeS-Ph	H	Me
1-182	H	NH(EtOCO)	4-MeS-Ph	H	Me
1-183	H	NH(BzOCO)	4-MeS-Ph	H	Me
1-184	H	NH(AcOCH ₂)	4-MeS-Ph	H	Me
1-185	H	NH(PmOCH ₂)	4-MeS-Ph	H	Me
1-186	H	NH(MeOCOOCH ₂)	4-MeS-Ph	H	Me
1-187	H	NH(EtOCOOCH ₂)	4-MeS-Ph	H	Me

Table 1 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
1-188	H	NH[(5-Me-2-oxo-1,3-dioxolen-4-yl)CH ₂]	4-MeS-Ph	H	Me
1-189	H	NH[(5-Ph-2-oxo-1,3-dioxolen-4-yl)CH ₂]	4-MeS-Ph	H	Me

Table 2

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
2-1	H	Me	Ph	H	H
2-2	H	Me	Ph	H	Me
2-3	H	Me	4-F-Ph	H	H
2-4	H	Me	4-F-Ph	F	H
2-5	H	Me	4-F-Ph	Cl	H
2-6	H	Me	4-F-Ph	Br	H
2-7	H	Me	4-F-Ph	I	H
2-8	H	Me	4-F-Ph	Me	H
2-9	H	Me	4-F-Ph	Et	H
2-10	H	Me	4-F-Ph	Pr	H
2-11	H	Me	4-F-Ph	H	Me
2-12	H	Me	4-F-Ph	H	Et
2-13	H	Me	4-F-Ph	H	Pr
2-14	H	Me	4-F-Ph	H	Bu
2-15	H	Me	4-F-Ph	H	cPr
2-16	H	Me	4-F-Ph	H	Ph
2-17	H	Me	4-F-Ph	H	CH ₂ Ph
2-18	H	Me	4-F-Ph	H	CHF ₂
2-19	H	Me	4-F-Ph	H	CF ₃
2-20	H	Me	4-MeO-Ph	H	H

Table 2 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
2-21	H	Me	4-MeO-Ph	Me	H
2-22	H	Me	4-MeO-Ph	H	Me
2-23	H	Me	4-Cl-Ph	H	H
2-24	H	Me	4-Cl-Ph	Me	H
2-25	H	Me	4-Me-Ph	H	H
2-26	H	Me	4-Me-Ph	Me	H
2-27	H	Me	4-Me-Ph	H	Me
2-28	H	Me	3-Cl-4-F-Ph	H	H
2-29	H	Me	3-Cl-4-F-Ph	H	Me
2-30	H	Me	3,4-Methylenedioxy-Ph	H	H
2-31	H	Me	3,4-Methylenedioxy-Ph	H	Me
2-32	H	Me	3-Cl-4-MeO-Ph	H	H
2-33	H	Me	3-Cl-4-MeO-Ph	H	Me
2-34	H	Me	4-CF ₃ -Ph	H	H
2-35	H	Me	4-CF ₃ O-Ph	H	H
2-36	H	Me	4-CHF ₂ O-Ph	H	H
2-37	H	Me	4-CHF ₂ O-Ph	Me	H
2-38	H	Me	3-F-4-MeO-Ph	H	H
2-39	H	Me	3-F-4-MeO-Ph	H	Me
2-40	H	Me	3-Me-4-MeO-Ph	H	H

Table 2 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
2-41	H	Me	3-Me-4-MeO-Ph	H	Me
2-42	H	Me	3,4-diF-Ph	H	H
2-43	H	Me	3,4-diF-Ph	H	Me
2-44	H	Me	2,4-diF-Ph	H	H
2-45	H	Me	2,4-diF-Ph	H	Me
2-46	H	Me	3,4-diMe-Ph	H	H
2-47	H	Me	3,4-diCl-Ph	H	H
2-48	H	Me	3,4-diCl-Ph	H	Me
2-49	H	Me	3,4-di(MeO)-Ph	H	H
2-50	H	Me	3,4-di(MeO)-Ph	H	Me
2-51	H	Me	4-F-Ph	H	CH ₂ OH
2-52	H	Me	4-F-Ph	H	CH ₂ OMe
2-53	H	Me	4-MeO-Ph	H	CH ₂ OH
2-54	H	Me	4-MeO-Ph	H	CH ₂ OMe
2-55	H	Me	4-Cl-Ph	H	CH ₂ OH
2-56	H	Me	4-Cl-Ph	H	CH ₂ OMe
2-57	H	Me	4-Me-Ph	H	CH ₂ OH
2-58	H	Me	4-Me-Ph	H	CH ₂ OMe
2-59	H	NH ₂	Ph	H	H

Table 2 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
2-60	H	NH ₂	Ph	H	Me
2-61	H	NH ₂	Ph	Me	H
2-62	H	NH ₂	4-F-Ph	H	H
2-63	H	NH ₂	4-F-Ph	H	Me
2-64	H	NH ₂	4-F-Ph	Me	H
2-65	H	NH ₂	4-F-Ph	H	Et
2-66	H	NH ₂	4-F-Ph	H	Pr
2-67	H	NH ₂	4-F-Ph	H	Bu
2-68	H	NH ₂	4-F-Ph	H	cPr
2-69	H	NH ₂	4-F-Ph	H	Ph
2-70	H	NH ₂	4-F-Ph	H	CH ₂ Ph
2-71	H	NH ₂	4-F-Ph	H	CHF ₂
2-72	H	NH ₂	4-F-Ph	H	CF ₃
2-73	H	NH ₂	4-MeO-Ph	H	H
2-74	H	NH ₂	4-MeO-Ph	H	Me
2-75	H	NH ₂	4-MeO-Ph	H	Et
2-76	H	NH ₂	4-MeO-Ph	Me	H
2-77	H	NH ₂	4-EtO-Ph	H	H

Table 2 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
2-96	H	NH ₂	3-Cl-4-F-Ph	Me	H
2-97	H	NH ₂	3,4-Methylenedioxy-Ph	H	H
2-98	H	NH ₂	3,4-Methylenedioxy-Ph	H	Me
2-99	H	NH ₂	3-Cl-4-MeO-Ph	H	H
2-100	H	NH ₂	3-Cl-4-MeO-Ph	H	Me
2-101	H	NH ₂	3-Cl-4-MeO-Ph	Me	H
2-102	H	NH ₂	4-CF ₃ -Ph	H	Me
2-103	H	NH ₂	4-CHF ₂ O-Ph	H	Me
2-104	H	NH ₂	4-CF ₃ O-Ph	H	Me
2-105	H	NH ₂	2-F-4-MeO-Ph	H	Me
2-106	H	NH ₂	3-F-4-MeO-Ph	H	Me
2-107	H	NH ₂	3-F-4-MeO-Ph	Me	H
2-108	H	NH ₂	3-Me-4-MeO-Ph	H	H
2-109	H	NH ₂	3-Me-4-MeO-Ph	H	Me
2-110	H	NH ₂	3-Me-4-MeO-Ph	Me	H
2-111	H	NH ₂	3,4-diF-Ph	H	H
2-112	H	NH ₂	3,4-diF-Ph	H	Me
2-113	H	NH ₂	3,4-diF-Ph	Me	H

Table 2 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
2-114	H	NH ₂	2,4-diF-Ph	H	H
2-115	H	NH ₂	2,4-diF-Ph	H	Me
2-116	H	NH ₂	2,4-diF-Ph	Me	H
2-117	H	NH ₂	3,4-diMe-Ph	H	H
2-118	H	NH ₂	3,4-diMe-Ph	H	Me
2-119	H	NH ₂	3,4-diMe-Ph	Me	H
2-120	H	NH ₂	2,4-diCl-Ph	H	H
2-121	H	NH ₂	2,4-diCl-Ph	H	Me
2-122	H	NH ₂	2,4-diCl-Ph	Me	H
2-123	H	NH ₂	3,4-diCl-Ph	H	H
2-124	H	NH ₂	3,4-diCl-Ph	H	Me
2-125	H	NH ₂	3,4-diCl-Ph	Me	H
2-126	H	NH ₂	3,4-di(MeO)-Ph	H	H
2-127	H	NH ₂	3,4-di(MeO)-Ph	H	Me
2-128	H	NH ₂	4-F-Ph	H	CH ₂ OH
2-129	H	NH ₂	4-F-Ph	H	CH ₂ OMe
2-130	H	NH ₂	4-MeO-Ph	H	CH ₂ OH
2-131	H	NH ₂	4-MeO-Ph	H	CH ₂ OMe

Table 2 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
2-132	H	NH ₂	4-Cl-Ph	H	CH ₂ OH
2-133	H	NH ₂	4-Cl-Ph	H	CH ₂ OMe
2-134	H	NH ₂	4-Me-Ph	H	CH ₂ OH
2-135	H	NH ₂	4-Me-Ph	H	CH ₂ OMe
2-136	H	NH ₂	3,5-diCl-4-MeO-Ph	H	Me
2-137	H	NH ₂	3,5-diMe-4-MeO-Ph	H	Me
2-138	H	NH ₂	2,3-diCl-Ph	H	Me
2-139	H	NH ₂	3,5-diCl-Ph	H	Me
2-140	H	NH ₂	2,4,5-triMe-Ph	H	Me
2-141	H	NH ₂	3-cPnO-4-MeO-Ph	H	Me
2-142	H	NH ₂	3-CF ₃ -4-Cl-Ph	H	Me
2-143	H	NH ₂	3-F-4-Me-Ph	H	Me
2-144	H	NH ₂	3-Me-4-Cl-Ph	H	Me
2-145	H	NH ₂	2,4-diMe-Ph	H	Me
2-146	H	NH ₂	4-OH-Ph	H	Me
2-147	H	NH ₂	3,5-diMe-Ph	H	Me
2-148	H	NHAc	4-MeO-Ph	H	Me
2-149	H	NHAc	3,4-diMe-Ph	H	Me

Table 2 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
2-150	H	NH ₂	4-MeO-Ph	H	3-cPnO-4-MeO-Bz
2-151	H	NH ₂	4-MeSO-Ph	H	Me
2-152	3-F	NH ₂	4-MeO-Ph	H	Me
2-153	3-F	NH ₂	4-EtO-Ph	H	Me
2-154	3-F	NH ₂	3,4-diMe-Ph	H	Me
2-155	3-F	NH ₂	4-Cl-Ph	H	Me
2-156	3-F	NH ₂	4-F-Ph	H	Me
2-157	3-F	NH ₂	4-SH-Ph	H	Me
2-158	3-F	NH ₂	4-MeS-Ph	H	Me
2-159	3-F	NH ₂	4-EtS-Ph	H	Me
2-160	3-F	NH ₂	4-AcS-Ph	H	Me
2-161	3-Me	NH ₂	4-MeO-Ph	H	Me
2-162	3-Me	NH ₂	4-EtO-Ph	H	Me
2-163	3-Me	NH ₂	3,4-diMe-Ph	H	Me
2-164	3-Me	NH ₂	4-MeS-Ph	H	Me
2-165	H	NHFor	4-MeO-Ph	H	Me
2-166	H	NHPm	4-MeO-Ph	H	Me
2-167	H	NHByr	4-MeO-Ph	H	Me

Table 2 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
2-168	H	NHiByr	4-MeO-Ph	H	Me
2-169	H	NHVal	4-MeO-Ph	H	Me
2-170	H	NHiVal	4-MeO-Ph	H	Me
2-171	H	NHPiv	4-MeO-Ph	H	Me
2-172	H	NH(MeOCO)	4-MeO-Ph	H	Me
2-173	H	NH(EtOCO)	4-MeO-Ph	H	Me
2-174	H	NH(BzOCO)	4-MeO-Ph	H	Me
2-175	H	NH(AcOCH ₂)	4-MeO-Ph	H	Me
2-176	H	NH(PmOCH ₂)	4-MeO-Ph	H	Me
2-177	H	NH(MeOCOCH ₂)	4-MeO-Ph	H	Me
2-178	H	NH(EtOCOCH ₂)	4-MeO-Ph	H	Me
2-179	H	NH[(5-Me-2-oxo-1,3-dioxolen-4-yl)CH ₂]	4-MeO-Ph	H	Me
2-180	H	NH[(5-Ph-2-oxo-1,3-dioxolen-4-yl)CH ₂]	4-MeO-Ph	H	Me
2-181	H	NHFor	4-EtO-Ph	H	Me
2-182	H	NHAc	4-EtO-Ph	H	Me
2-183	H	NHPm	4-EtO-Ph	H	Me
2-184	H	NHByr	4-EtO-Ph	H	Me
2-185	H	NHiByr	4-EtO-Ph	H	Me

Table 2 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
2-186	H	NHVal	4-EtO-Ph	H	Me
2-187	H	NHiVal	4-EtO-Ph	H	Me
2-188	H	NHPiv	4-EtO-Ph	H	Me
2-189	H	NH(MeOCO)	4-EtO-Ph	H	Me
2-190	H	NH(EtOCO)	4-EtO-Ph	H	Me
2-191	H	NH(BzOCO)	4-EtO-Ph	H	Me
2-192	H	NH(AcOCH ₂)	4-EtO-Ph	H	Me
2-193	H	NH(PmOCH ₂)	4-EtO-Ph	H	Me
2-194	H	NH(MeOCOOCH ₂)	4-EtO-Ph	H	Me
2-195	H	NH(EtOCOOCH ₂)	4-EtO-Ph	H	Me
2-196	H	NH[(5-Me-2-oxo-1,3-dioxolen-4-yl)CH ₂]	4-EtO-Ph	H	Me
2-197	H	NH[(5-Ph-2-oxo-1,3-dioxolen-4-yl)CH ₂]	4-EtO-Ph	H	Me
2-198	H	NHFor	3,4-diMe-Ph	H	Me
2-199	H	NHPm	3,4-diMe-Ph	H	Me
2-200	H	NHByr	3,4-diMe-Ph	H	Me
2-201	H	NHiByr	3,4-diMe-Ph	H	Me
2-202	H	NHVal	3,4-diMe-Ph	H	Me
2-203	H	NHiVal	3,4-diMe-Ph	H	Me

Table 2 (cont.)

Cpd. No.	R	R ¹	R ²	R ³	R ⁴
2-204	H	NHPiv	3,4-diMe-Ph	H	Me
2-205	H	NH(MeOCO)	3,4-diMe-Ph	H	Me
2-206	H	NH(EtOCO)	3,4-diMe-Ph	H	Me
2-207	H	NH(BzOCO)	3,4-diMe-Ph	H	Me
2-208	H	NH(AcOCH ₂)	3,4-diMe-Ph	H	Me
2-209	H	NH(PmOCH ₂)	3,4-diMe-Ph	H	Me
2-210	H	NH(MeOCOOCH ₂)	3,4-diMe-Ph	H	Me
2-211	H	NH(EtOCOOCH ₂)	3,4-diMe-Ph	H	Me
2-212	H	NH[(5-Me-2-oxo-1,3-dioxolen-4-yl)CH ₂]	3,4-diMe-Ph	H	Me
2-213	H	NH[(5-Ph-2-oxo-1,3-dioxolen-4-yl)CH ₂]	3,4-diMe-Ph	H	Me

Of the compounds listed above, preferred compounds are:

- 1) 3-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,
- 2) 4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,
- 3) 1-(4-fluorophenyl)-2-(4-sulfamoylphenyl)pyrrole,
- 4) 1-(4-fluorophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
- 5) 5-fluoro-1-(4-fluorophenyl)-2-(4-methylsulfonylphenyl)pyrrole,
- 6) 2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

- 7) 1-(4-methoxyphenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
- 8) 4-ethyl-2-(4-methoxyphenyl)-1-(4-sulfamoylphenyl)pyrrole,
- 9) 2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 10) 4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl)pyrrole,
- 5 11) 2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 12) 2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 13) 2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 14) 4-methyl-2-phenyl-1-(4-sulfamoylphenyl)pyrrole,
- 15) 2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 10 16) 2-(3-chloro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 17) 4-methyl-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole,
- 18) 5-chloro-1-(4-methoxyphenyl)-2-(4-sulfamoylphenyl)pyrrole,
- 19) 1-(3,4-dimethylphenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
- 20) 5-chloro-1-(4-ethoxyphenyl)-2-(4-sulfamoylphenyl)pyrrole,
- 15 21) 5-chloro-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole,
- 22) 1-(4-ethylthiophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
- 23) 2-(3,5-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 24) 1-(4-mercaptophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
- 25) 1-(4-acetylthiophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
- 20 26) 1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(4-methoxyphenyl)pyrrole, and
- 27) 1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(3,4-dimethylphenyl)pyrrole.

Of these, more preferred compounds are:

- 2) 4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,
- 6) 2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 9) 2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 5 10) 4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl)pyrrole,
- 11) 2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 12) 2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 13) 2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 15) 2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 10 17) 4-methyl-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole,
- 26) 1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(4-methoxyphenyl)pyrrole, and
- 27) 1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(3,4-dimethylphenyl)pyrrole.

Of these, the most preferred compounds are:

- 11) 2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 15 15) 2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
- 17) 4-methyl-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole,
- 26) 1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(4-methoxyphenyl)pyrrole, and
- 27) 1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(3,4-dimethylphenyl)pyrrole.

The compounds of formula (I), compounds of formula (II) and
20 pharmaceutically acceptable salts of these compounds are known compounds and a method of preparing these compounds is disclosed in European Patent Publication EP-799823A, the disclosure of which is incorporated herein by reference.

The chemical names of the compounds of formulae (III) to (XIV), respectively, are:

- (III): 3-(3,4-difluorophenyl)-4-(4-methanesulfonylphenyl)-5H-furan-2-one,
- (IV): 4-(5-p-tolyl-3-trifluoromethyl-1H-pyrazol-1-yl)benzenesulfonamide,
- 5 (V): N-[6-(2,4-difluorophenylthio)-1-oxoindan-5-yl]methanesulfonamide,
- (VI): 4-hydroxy-2-methyl-N-(5-methylthiazol-2-yl)-2H-1,2-benzothiazine-3-carboxamide-1,1-dioxide,
- (VII): N-(4-Nitro-2-phenoxyphenyl)methanesulfonamide,
- (VIII): 4-(4-cyclohexyl-2-methyloxazol-5-yl)-2-fluorobenzenesulfonamide,
- 10 (IX): N-(3-formylamino-4-oxo-6-phenoxy-4H-1-benzopyran-7-yl)methanesulfonamide,
- (X): (E)-2-ethyl-5-(3,5-di-t-butyl-4-hydroxy)benzylidene-1,2-isothiazolidine-1,1-dioxide,
- (XI): 1-(4-methanesulfonylphenyl)-2-(4-fluorophenyl)cyclopentene,
- 15 (XII): 3-phenyl-4-(4-methanesulfonylphenyl)-5H-furan-2-one, and
- (XIII): 2-(3,5-difluorophenyl)-3-(4-methanesulfonylphenyl)-2-cyclopenten-1-one.
- (XIV): 4-[5-methyl-3-phenylisoxazol-4-yl]benzenesulfonamide.

These compounds are disclosed in International publication number

20 WO95/00501, J. Med. Chem., 40, 1347 (1997), International publication number WO94/13635, Pharmacology, 55, 44 (1997), Prostaglandins, 47, 55 (1994), Japanese publication number Hei 9-52882, Jpn. J. Pharmacol., 67, 305 (1995), Inflamm. Res., 47, Suppl. 3, S257 (1997), J. Med. Chem., 38, 4570 (1995), EP 863 134, US 5 474 995 or WO 98/06708, the disclosures of which are incorporated herein by

25 reference.

Since the compounds of the present invention have excellent activity for the prevention or inhibition of cachexia and very little toxicity, they are useful as preventive and therapeutic agents for cachexia. They are also useful for the treatment of tumor-related disorders, and can be used to inhibit the growth and/or metastasis of
5 tumors.

Moreover, if desired, one or more of the compounds of the present invention [i.e. the compounds of formulae (I) to (XIV), inclusive] may be used in association with one or more other agents for the prevention or inhibition for tumor growth, and the compounds of the present invention and other agents may be administered
10 simultaneously, separately or sequentially.

The other antitumor agent is preferably selected from a group consisting of 5-fluorouracil, cisplatin, tamoxifen, paclitaxel, docetaxel and irinotecan. Especially in the case of simultaneous administration, the compound of formula (I) or (II) or a pharmaceutically acceptable salt thereof and the other antitumor agent may be
15 contained in a single composition.

The composition of the present invention may be in any conventional form, depending on the route of administration. For example, for oral administration, it may be in the form of tablets, capsules, granules, powders or syrups. For non-oral administration it may be in the form of injections or suppositories. These
20 formulations are prepared according to known methods and may include additives such as are well known in the art, for example excipients (e.g., organic excipients including sugar derivatives, such as lactose, sucrose, glucose, mannitol and sorbitol; starch derivatives, such as corn starch, potato starch, α -starch and dextrin; cellulose derivatives, such as crystalline cellulose; gum arabic; dextran; and Pullulan,
25 inorganic excipients including silicate derivatives, such as light silicic acid anhydride, synthetic aluminum silicate, calcium silicate and magnesium metasilicate aluminate; phosphates, such as calcium hydrogenphosphate; carbonates, such as calcium carbonate; and sulfates, such as calcium sulfate), lubricants (e.g., stearic acid and metal salts thereof, including stearic acid, calcium stearate and magnesium
30 stearate; talc; colloidal silica; waxes, such as beeswax and spermaceti; boric acid;

adipic acid; sulfates, such as sodium sulfate; glycol; fumaric acid; sodium benzoate; DL-leucine; fatty acid sodium salts; lauryl sulfates, such as sodium lauryl sulfate and magnesium lauryl sulfate; silicic acids, such as silicic acid anhydride and silicic acid hydrate; and the above-mentioned starch derivatives), binders (e.g., hydroxypropyl
5 cellulose, hydroxypropylmethyl cellulose, polyvinyl pyrrolidone, Macrogol and similar compounds to the above-mentioned excipients), disintegrating agents (e.g., cellulose derivatives, such as low-substituted hydroxypropyl cellulose, carboxymethyl cellulose, calcium carboxymethyl cellulose, internally bridged sodium carboxymethyl cellulose; chemically modified starch-celluloses, such as
10 carboxymethyl starch, sodium carboxymethyl starch and bridged polyvinyl pyrrolidone), stabilizers (e.g., paraoxybenzoates, such as methylparaben and propylparaben; alcohols, such as chlorobutanol, benzyl alcohol and phenylethyl alcohol; benzalkonium chloride; phenols, such as phenol and cresol; thimerosal; dehydroacetic acid; and sorbic acid), corrigents (e.g., sweeteners, vinegars and
15 perfumes) and diluents.

The dose varies, depending on many factors, including the condition and age of the patients, the severity and nature of the disorder and the route of administration. For example, in the case of oral administration, it is desirable to administer 0.01 mg/kg (preferably 0.1 mg/kg) as a lower limit and 50 mg/kg (preferably 10 mg/kg) as
20 an upper limit for an adult per day, in a single dose or in divided doses, depending on the symptoms. In the case of intravenous administration, it is desirable to administer 0.001 mg/kg (preferably 0.01 mg/kg) as a lower limit and 10 mg/kg (preferably 5 mg/kg) as an upper limit for an adult, in a single dose or in divided doses, depending on the symptoms.

25 The present invention is further illustrated by the following non-limiting Examples and Formulation examples.

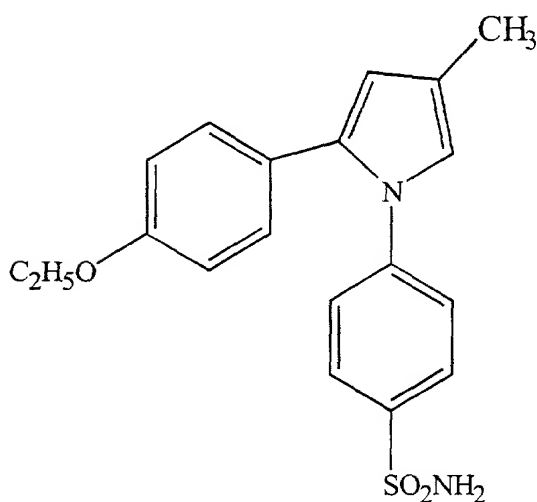
EXAMPLE 1

Test of Anticachexia Effects in Mice bearing Mouse Colon Cancer Colon 26

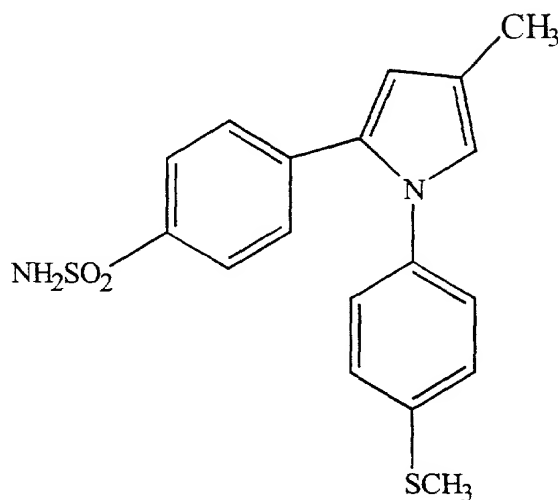
Cells

The test animals were CDF1 mice (females, 8 weeks old). They were employed
5 in groups of 10 for each test. 1×10^6 mouse colon cancer Colon 26 cells were
transplanted subcutaneously into each animal.

The test compounds were Compounds No. 1-94 and 2-78 as shown above in
Tables 1 and 2, respectively, and having the following formulae:



Compound No. 2-78



Compound No. 1-94

10

Each test compound was suspended in sterilized distilled water containing
0.5% w/v carboxymethyl cellulose (CMC) and administered orally once per day
starting on the day of the tumor cell transplantation.

Each test animal was weighed immediately after tumor cell transplantation, and
15 the weight (A g) was recorded. Each animal was then weighed on day 19 after tumor
cell transplantation, and the weight (B g) was recorded. The weight gain on day 19
after tumor cell transplantation was calculated as $B - A = \Delta g_t$ for the test animals. The
experiment was repeated with two control groups: the first control group (control

group 1) was transplanted with the tumor cells but were not treated with any test compound, and the weight gain is reported as Δg_{c1} ; the second control group (control group 2) were not transplanted with the tumor cells and were not treated with any test compound, and the weight gain is reported as Δg_{c2} . The body weight recovery rate was determined according to the following formula based on the weight gain on day 19 after tumor cell transplantation, and this value was used as an indicator of the anticachexia effect.

$$\text{Body weight recovery rate (\%)} = (\Delta g_t - \Delta g_{c1}) / (\Delta g_{c2} - \Delta g_{c1}) \times 100$$

The results are shown in Table 3 below.

Table 3

Administered Compound	Dose (mg/kg)	Weight Gain (Δg)	Body Weight Recovery Rate (%)
Compound 2-78	10	2.6	88
Compound 2-78	3	2.5	85
Compound 2-78	1	2.5	85
Compound 1-94	10	3.1	98
Compound 1-94	3	2.7	89
Compound 1-94	1	2.2	78
control group 1	-	-1.4	0
control group 2	-	3.2	100

It is clear from the above results that these compounds inhibited mouse tumor cachexia and reduced weight loss.

EXAMPLE 2

Test of Anticachexia Effects in Mice bearing Mouse Colon Cancer Colon 26 Cells

The procedure described in Example 1 was repeated, but using the compound of formula (III) as the test compound, and comparing the weight gain (Δg_t) of the test group of animals to which the compound of formula (III) had been administered with a control group (Δg_c) into which the tumor cells had been transplanted but to which no anti-tumor compound had been administered. The test animals were female CDF1 mice, 16 weeks old. Also, the weight gain was measured 22 days after tumor transplantation. The average body weight of each group of animals immediately after tumor transplantation was 25 to 26 g. The results are shown in Table 4.

Table 4

Compound	Dose (mg/kg)	Average Weight Gain (Δg)
Compound (III)	10	0.9
Compound (III)	3	0.3
Compound (III)	1	0.0
None (Control group)	-	-4.2

It is clear from the above results that the compound of formula (III) inhibited mouse tumor cachexia and reduced weight loss.

EXAMPLE 3

Test of Anticachexia Effects in Mice bearing Mouse Colon Cancer Colon 26 Cells

The procedure described in Example 2 was repeated, but using the compound of formula (IV) as the test compound, and comparing the weight gain (Δg_t) of the test

group of animals to which the compound of formula (IV) had been administered with a control group (Δg_c) into which the tumor cells had been transplanted but to which no anti-tumor compound had been administered. The test animals were female CDF1 mice, 7 weeks old. Also, the weight gain was measured 15 days after tumor transplantation. The average body weight of each group of animals immediately after tumor transplantation was 20 to 21 g. The results are shown in Table 5.

Table 5

Compound	Dose (mg/kg)	Average Weight Gain (Δg)
Compound (IV)	10	-0.6
Compound (IV)	3	-1.3
Compound (IV)	1	-1.2
None (Control group)	-	-3.4

It is clear from the above results that the compound of formula (IV) inhibited mouse tumor cachexia and reduced weight loss.

EXAMPLE 4

Test of Anticachexia Effects in Mice bearing Mouse Colon Cancer Colon 26 Cells

Test compounds [the compounds of formula (V), (VI), (VII), (VIII), (IX), (X), (XI), (XII), (XIII) and (XIV)] are administered in the same manner as described in Example 2. These compounds inhibit mouse tumor cachexia and recover loss of average of body weight.

EXAMPLE 5

Life-prolonging Activity Test

Observation of the mice used in Example 1 above was continued. The life-prolonging index was determined, based on the number of days each mouse survived, and this value was then used as an indicator of the life-prolonging effects of the test

compounds.

It should be noted that, in the case of the group of mice treated with a test compound, oral administration of the respective compound once daily was continued on day 20 after tumor cell transplantation and beyond as well.

5
$$\text{Life-prolonging index (\%)} = (S_t/S_c - 1) \times 100$$

S_t : Median value of survival time (days) of the group of mice treated with a test compound

S_c : Median value of survival time (days) of the control group which were not transplanted with tumor cells.

10 The results are shown in Table 6.

Table 6

Compound Name	Dose (mg/kg)	Survival Period (median: days)	Life-prolonging Index (%)
Compound 2-78	10	48.5	73
Compound 2-78	3	50.5	80
Compound 2-78	1	45.0	61
Compound 1-94	10	45.0	61
Compound 1-94	3	35.0	25
Compound 1-94	1	48.5	73
None	-	28.0	0

As is clear from Table 6, the compounds of the present invention exhibited a prominent life-prolonging effect.

EXAMPLE 6

Life-prolonging Activity Test

The experiment reported in Example 5 was repeated with the animals used in Example 2. In the case of the group of mice treated with a test compound, oral
5 administration of the respective compound once daily was continued on day 23 after tumor cell transplantation and beyond as well. The results are shown in Table 7.

Table 7

Compound Name	Dose (mg/kg)	Survival Period (median: days)	Life-prolonging Index (%)
Compound (III)	10	43.5	91
Compound (III)	3	37.5	63
Compound (III)	1	40.5	76
None	-	23.0	-

EXAMPLE 7

Life-prolonging Activity Test

The experiment reported in Example 5 is repeated with the animals used in Examples 3 and 4. The compounds of formulae (IV), (V), (VI), (VII), (VIII), (IX), (X), (XI), (XII), (XIII) and (XIV) all inhibit mouse tumor cachexia and exhibit pronounced life-prolongation.

EXAMPLE 8

Test of concomitant use of antitumor agent

15 Mouse colon cancer cells are transplanted into CDF1 mice in the same manner as Example 1 followed by administration of the test compounds [Compound Nos. 2-78 and 1-94, and the compounds of formulae (III), (IV), (V), (VI), (VII), (VIII), (IX),
20 (X), (XI), (XII), (XIII) and (XIV)] and an antitumor agent (5-fluorouracil or cisplatin).

[illegible]

5 **Inhibitory effect on lung metastasis of mouse malignant melanoma B16-BL6**
cells

The mice were, when necessary, administered intravenously into the tail vein with a bacterial lipopolysaccharide (LPS) in an amount of 3 μ g each within one hour before transplantation of the melanoma cells so as to accelerate lung metastasis of the melanoma [M. J. Anasagasti et al., J. Natl. Cancer Research, 89, 645-651 (1997).]

Compound No. 2-118 is 2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoyl-
20 phenyl)pyrrole.

$$25 \quad \text{LMI (\%)} = (1 - N_t / N_c) \times 100$$

N_t : Numbers of lung metastatic colonies on the 10th day in groups administered with the test compounds; and

N_C: Numbers of lung metastatic colonies on the 10th day in control groups which were not administered with the test compounds.

The results are shown in Table 8.

Table 8

Test compound	LPS administration	Dose (mg/kg)	LMI (%)
Compound 2-118	No	1	64
Compound 2-118	Yes	1	34
Compound (IV)	Yes	1	9
Compound (V)	Yes	1	1
Indomethacin	Yes	1	-1

5

It is clear from Table 8 that the present composition was successful in inhibiting metastasis of the mouse malignant melanoma B16-BL6 cells to the lung whether or not lung metastasis was accelerated by the LPS administration (induction of inflammation reaction).

10 In particular, the present composition showed a marked inhibition of lung metastasis, while the compound of formula (IV) and the compound of formula (V), which are COX-2-selective inhibitors, and indomethacin, which is a typical NSAID, had no such inhibitory activity under the lung metastasis accelerating conditions caused by inducing inflammatory reaction (as reflecting acceleration of metastasis of
15 tumor in a surgical operation of resecting a tumor).

EXAMPLE 10

Antitumor Effect Against Mouse Sarcoma S-180 Cells

1 x 10⁶ mouse sarcoma S-180 cells were transplanted subcutaneously in Balb/c nude mice (females, 8 weeks old) in groups of 10 each.

The test compound, Compound No. 2-118, was suspended in sterilized distilled water containing 0.5% w/v carboxymethyl cellulose (CMC) and administered orally once per day for 5 days starting on the day the tumor cells were transplanted.

- 5 Antitumor activity was assessed according to the following equation to determine the tumor growth inhibitory rate (GI%) on day 7 after the transplantation.

$$GI (\%) = (1 - V_t / V_c) \times 100$$

V_t : Mean tumor volume on day 7 in a group administered test compound (*)

V_c : Mean tumor volume on day 7 in an untreated control group (*)

- 10 *: Tumor volume is defined as $1/2 \times [\text{tumor long axis}] \times [\text{tumor short axis}]^2$

The results are shown in Table 9.

Table 9

Test Compound	Dose (mg/kg)	GI (%)
Compound 2-118	1	54

- It is clear from Table 9 that the composition of the present application
15 inhibited the growth of mouse tumor cells.

EXAMPLE 11

Antitumor Effect Against Human Colon Cancer KM12-HX Cells

- Human colon cancer KM12-HX cells were orthotopically transplanted into the cecum of nude mice according to the method of Fu *et al.* [X. Fu *et al.*, Anticancer
20 Res., 12 (1992)] using Balb/c nude mice (females, 7 weeks old) in groups of 10 each.

Specifically, an incision was made into the left lower abdominal region of each mouse under Abacin anesthesia, after which a thin section of tumor measuring 5 mm

on a side was sutured to the cecum using absorbable surgical sutures to perform orthotopic transplant. The incision was sutured using absorbable surgical sutures and the mice were warmed and promptly awakened from anesthesia.

The test compound was suspended in sterilized distilled water containing 0.5% w/v carboxymethyl cellulose (CMC) and administered orally in a total of 9 doses consisting of one dose per day from days 3 to 7 after the tumor cell transplantation and from days 10 to 13 after the transplantation.

Antitumor activity was assessed according to the following equation to determine the tumor weight inhibitory rate (GI%) on day 14 after the transplantation.

$$GI (\%) = (1 - V_t' / V_c') \times 100$$

V_t' : Mean tumor weight on day 14 in the group administered the test compound

V_c' : Mean tumor weight on day 14 in an untreated control group

The results are shown in Table 10.

Table 10

Test Compound	Dose (mg/kg)	GI(%)
Compound 2-118	0.3	12
Compound 2-118	1	35
Compound 2-118	3	45
Compound 2-118	10	59

It is clear from Table 10 that the composition of the present invention inhibited the growth of human colon cancer cells at the orthotopic transplantation site.

Preparation of pharmaceutical formulations containing the compounds of the present invention is further illustrated by the following non-limiting Formulation Examples.

FORMULATION EXAMPLE 1

5 **Capsules**

A mixture of a compound of the present invention, such as the compound of formula (III), Compound No. 1-94, 2-78 or 2-118, is prepared in a digestive oily substance, such as soybean oil, cottonseed oil or olive oil, and filled into gelatin with a positive replacement pump to obtain soft capsules containing 100 mg of active
10 ingredient. The resulting capsules are then washed and dried.

FORMULATION EXAMPLE 2

Tablets

Tablets are manufactured in accordance with conventional methods using 100 mg of a compound of the present invention, such as the compound of formula
15 (III), Compound No. 1-94, 2-78 or 2-118, 0.2 mg of colloidal silicon dioxide, 5 mg of magnesium stearate, 275 mg of microcrystalline cellulose, 11 mg of starch and 98.8 mg of lactose.

In this case, the tablets can be coated with a preparation coating if desired.

FORMULATION EXAMPLE 3

20 **Injections**

1.5% by weight of a compound of the present invention, such as the compound of formula (III), Compound No. 1-94, 2-78 or 2-118, is stirred in 10% by volume of propylene glycol, and is then adjusted to a constant volume by the addition of water for injection, after which it was sterilized to prepare injections.

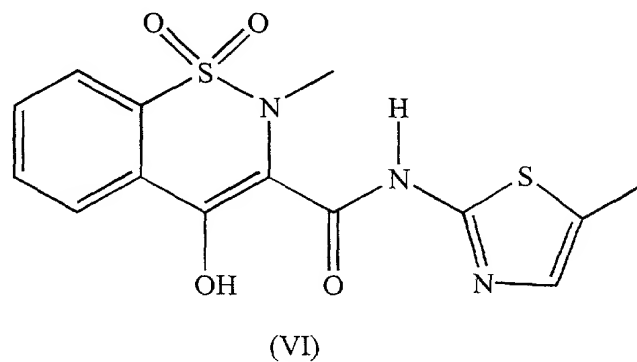
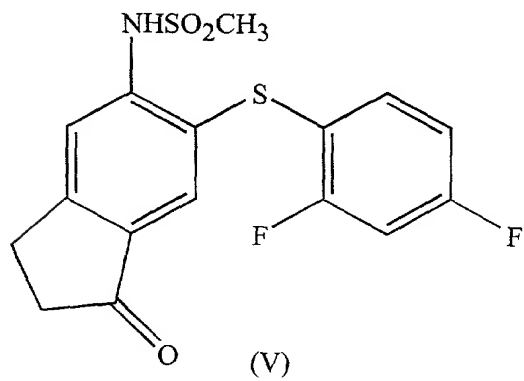
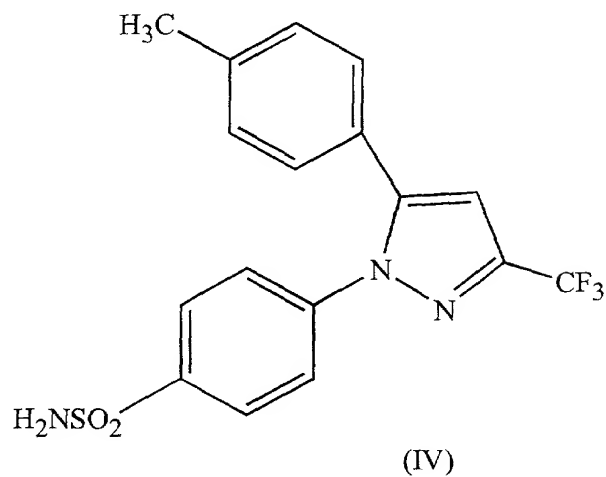
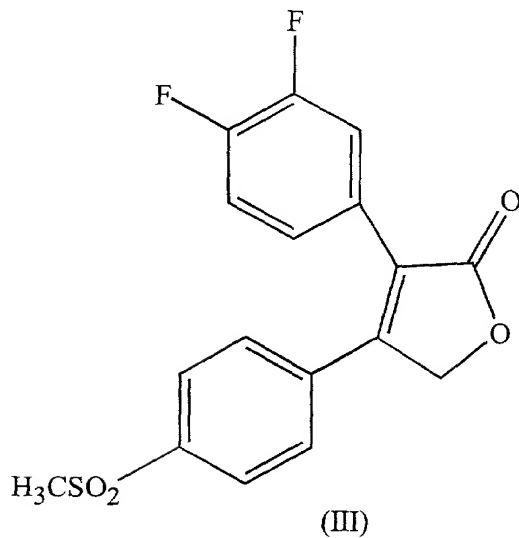
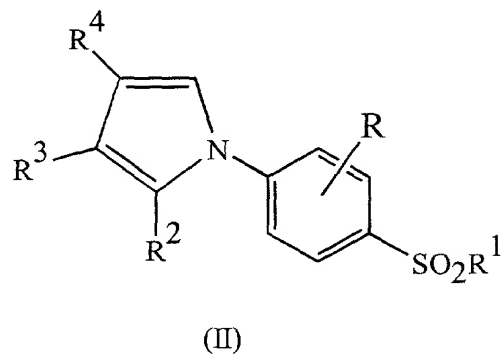
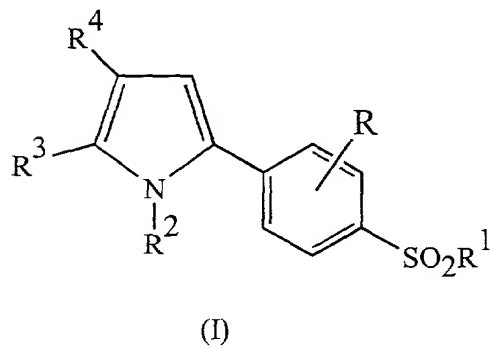
FORMULATION EXAMPLE 4

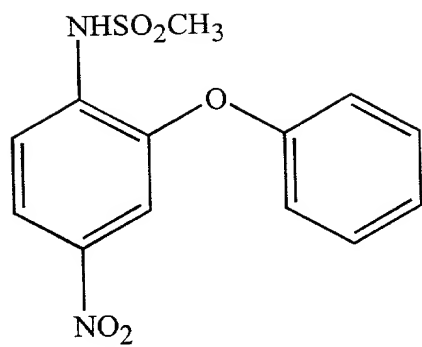
Suspensions

A suspension is produced so as to contain 100 mg of a compound of the present invention, such as the compound of formula (III), Compound No. 1-94, 2-78 or 2-
5 118, which is ground into a fine powder, 100 mg of sodium carboxymethyl cellulose, 5 mg of sodium benzoate, 1.0 g of sorbitol solution (Japanese Pharmacopoeia) and 0.025 ml of vanillin in 5 ml of the suspension.

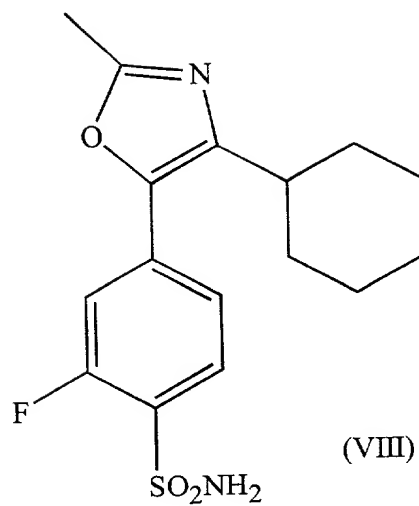
WE CLAIM:

1. A method for the treatment or prevention of cachexia in a mammal in need of such treatment or prevention, which method comprises administering to said mammal an effective amount of an active compound selected from the group consisting of compounds of formula (I), (II), (III), (IV), (V), (VI), (VII), (VIII), (IX), (X), (XI), (XII), (XIII) and (XIV):

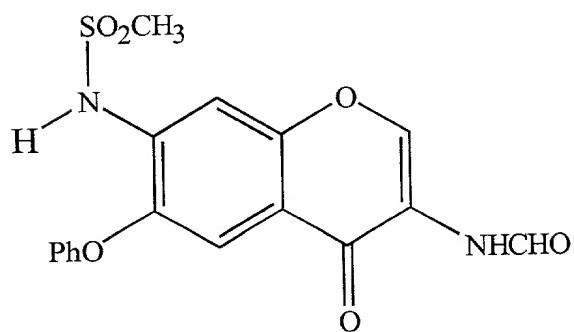




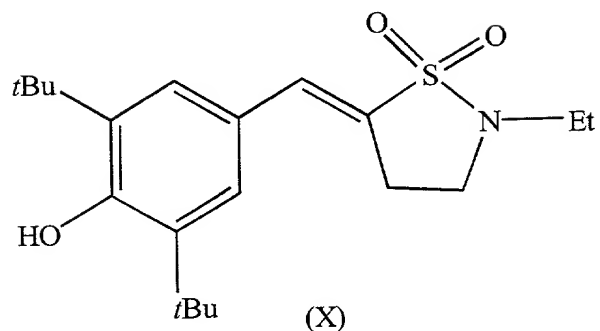
(VII)



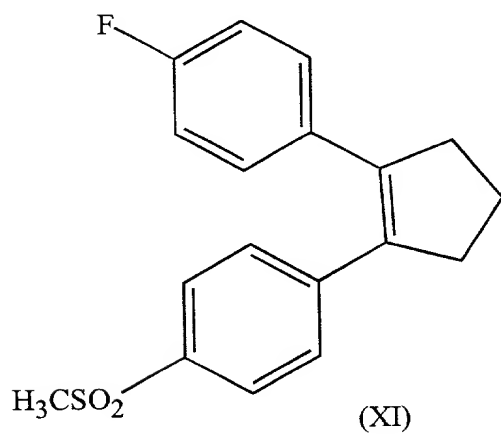
(VIII)



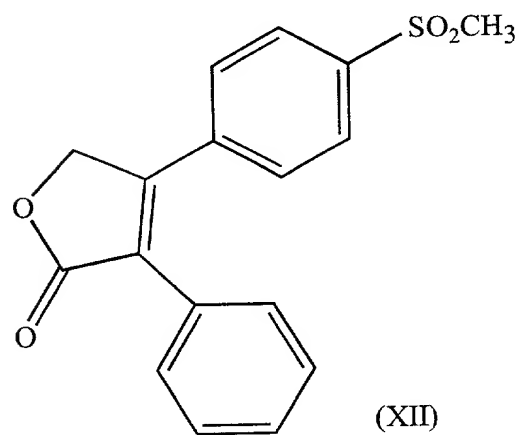
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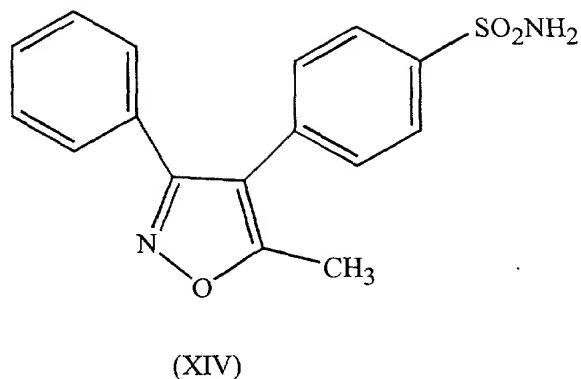
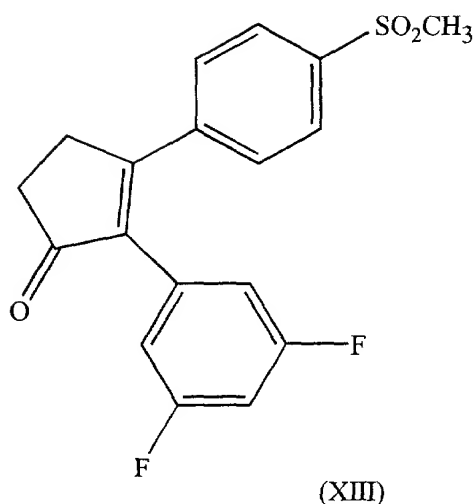
(X)



(XI)



(XII)



wherein

R represents a hydrogen atom, a halogen atom or a lower alkyl group;

R^1 represents a lower alkyl group, an amino group or a group of formula -

5 NHR^a (wherein R^a represents a group which may be eliminated *in vivo*);

R^2 represents a phenyl group or a phenyl group having at least one substituent selected from the group consisting of substituents α and substituents β , defined below;

10 R^3 represents a hydrogen atom, a halogen atom, a lower alkyl group or a lower alkyl group having at least one substituent selected from the group consisting of substituents α ;

15 R^4 represents a hydrogen atom, a lower alkyl group, a lower alkyl group having at least one substituent selected from the group consisting of substituents α , a cycloalkyl group, an aryl group as defined below, or an aralkyl group as defined below;

said aryl group is a carbocyclic aromatic hydrocarbon group having from 6 to 14 carbon atoms in one or more aromatic rings or such a group which is fused to a cycloalkyl group having from 3 to 10 carbon atoms, and the group is unsubstituted or it is substituted by at least one substituent selected from the

group consisting of substituents α and substituents β ;

said aralkyl group is a lower alkyl group which is substituted by one or more of the aryl groups defined above;

*t*Bu represents a t-butyl group;

5 Et represents an ethyl group; and

Ph represents a phenyl group;

said substituents α are selected from the group consisting of hydroxy groups, halogen atoms, lower alkoxy groups and lower alkylthio groups; and

10 said substituents β are selected from the group consisting of lower alkyl groups, alkanoyloxy groups, mercapto groups, alkanoylthio groups, lower alkylsulfinyl groups, lower alkyl groups having at least one substituent selected from the group consisting of substituents α , cycloalkyloxy groups, lower haloalkoxy groups and lower alkylenedioxy groups;

and pharmaceutically acceptable salts thereof.

- 15 2. The method of Claim 1, wherein said active compound is selected from the group consisting of compounds of formula (I) and (II).
3. The method of Claim 2, wherein R represents a hydrogen atom, a fluorine atom, a chlorine atom or a methyl group.
4. The method of Claim 2, wherein R represents a hydrogen atom.
- 20 5. The method of Claim 2, wherein R^1 represents a methyl group, an amino group or an acetylamino group.
6. The method of Claim 2, wherein R^1 represents an amino group or an acetylamino group.
7. The method of Claim 2, wherein R^2 represents a phenyl group or a phenyl group

substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^1 .

substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups; and

5 substituents β^1 are selected from the group consisting of lower alkyl groups, mercapto groups, alkanoylthio groups, lower alkyl groups substituted with at least one substituent selected from the group consisting of substituents α^1 , lower haloalkoxy groups and lower alkylenedioxy groups.

8. The method of Claim 2, wherein R^2 represents a phenyl group or a phenyl group
10 substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^2 .

substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups.

15 substituents β^2 are selected from the group consisting of lower alkyl groups, mercapto groups, alkanoylthio groups, lower alkyl groups substituted with a halogen atom, lower haloalkoxy groups and lower alkylenedioxy groups.

9. The method of Claim 2, wherein R^3 represents a hydrogen atom, a halogen atom, a lower alkyl group or a lower alkyl group substituted with at least one substituent selected from the group consisting of substituents α^1 .

20 substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups.

10. The method of Claim 2, wherein R^3 represents a hydrogen atom, a halogen atom, a lower alkyl group or a lower alkyl group substituted with a halogen atom.

11. The method of Claim 2, wherein R^4 represents a hydrogen atom, a lower alkyl
25 group, a lower alkyl group substituted with at least one substituent selected from the

group consisting of substituents α , a cycloalkyl group, an aryl group, an aryl group substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^3 , an aralkyl group or an aralkyl group substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^3 .

substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups; and

substituents β^3 are selected from the group consisting of lower alkyl groups, lower alkyl groups substituted with at least one substituent selected from the group consisting of substituents α and cycloalkyloxy groups.

12. The method of Claim 2, wherein R^4 represents a hydrogen atom, a lower alkyl group, a lower alkyl group substituted with at least one substituent selected from the group consisting of substituents α^2 , a cycloalkyl group, an aryl group, an aryl group substituted with at least one substituent selected from the group consisting of substituents α^2 and substituents β^4 , an aralkyl group or an aralkyl group substituted with at least one substituent selected from the group consisting of substituents α^2 and substituents β^4 .

substituents α^2 are selected from the group consisting of hydroxy groups, halogen atoms and lower alkoxy groups; and

substituents β^4 are selected from the group consisting of lower alkyl groups, lower alkyl groups substituted with halogen atom and cycloalkyloxy groups.

13. The method of Claim 2, wherein said active compound is:

3-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,

4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,

1-(4-fluorophenyl)-2-(4-sulfamoylphenyl)pyrrole,

- 1-(4-fluorophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
5-fluoro-1-(4-fluorophenyl)-2-(4-methylsulfonylphenyl)pyrrole,
2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
1-(4-methoxyphenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
5 4-ethyl-2-(4-methoxyphenyl)-1-(4-sulfamoylphenyl)pyrrole,
2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl)pyrrole,
2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
10 2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
4-methyl-2-phenyl-1-(4-sulfamoylphenyl)pyrrole,
2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
2-(3-chloro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
4-methyl-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole,
15 5-chloro-1-(4-methoxyphenyl)-2-(4-sulfamoylphenyl)pyrrole,
1-(3,4-dimethylphenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
5-chloro-1-(4-ethoxyphenyl)-2-(4-sulfamoylphenyl)pyrrole,
5-chloro-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole,
1-(4-ethylthiophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
20 2-(3,5-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
1-(4-mercaptophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,

1-(4-acetylthiophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,

1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(4-methoxyphenyl)pyrrole, or

1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(3,4-dimethylphenyl)pyrrole,

or a pharmaceutically acceptable salt thereof.

- 5 14. The method of Claim 2, wherein said active compound is:

2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

4-methyl-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole, or

- 10 1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(3,4-dimethylphenyl)pyrrole,

or a pharmaceutically acceptable salt thereof.

15. The method of Claim 1, wherein said active compound is selected from the group consisting of compounds of formulae (III), (IV), (V), (VI), (VII), (VIII), (IX), (X) and (XI).

- 15 16. The method of Claim 15, wherein said active compound is selected from the group consisting of 3-(3,4-difluorophenyl)-4-(4-methanesulfonylphenyl)-5H-furan-2-one and pharmaceutically acceptable salts thereof.

17. The method of Claim 15, wherein said active compound is selected from the group consisting of 4-(5-p-tolyl-3-trifluoromethyl-1H-pyrazol-1-yl)benzene-sulfonamide and pharmaceutically acceptable salts thereof.
- 20

18. The method of Claim 15, wherein said active compound is selected from the group consisting of N-[6-(2,4-difluorophenylthio)-1-oxoindan-5-yl]methane-sulfonamide and pharmaceutically acceptable salts thereof.

19. The method of Claim 15, wherein said active compound is selected from the

group consisting of 4-hydroxy-2-methyl-N-(5-methylthiazol-2-yl)-2H-1,2-benzothiazine-3-carboxamide-1,1-dioxide and pharmaceutically acceptable salts thereof.

20. The method of Claim 15, wherein said active compound is selected from the group consisting of N-(4-Nitro-2-phenoxyphenyl)methanesulfonamide and pharmaceutically acceptable salts thereof.

21. The method of Claim 15, wherein said active compound is selected from the group consisting of 4-(4-cyclohexyl-2-methyloxazol-5-yl)-2-fluorobenzene-sulfonamide and pharmaceutically acceptable salts thereof.

22. The method of Claim 15, wherein said active compound is selected from the group consisting of N-(3-formylamino-4-oxo-6-phenoxy-4H-1-benzopyran-7-yl)methanesulfonamide and pharmaceutically acceptable salts thereof.

23. The method of Claim 15, wherein said active compound is selected from the group consisting of (E)-2-ethyl-5-(3,5-di-t-butyl-4-hydroxy)benzylidene-1,2-isothiazolidine-1,1-dioxide and pharmaceutically acceptable salts thereof.

24. The method of Claim 15, wherein said active compound is selected from the group consisting of 1-(4-methanesulfonylphenyl)-2-(4-fluorophenyl)cyclopentene and pharmaceutically acceptable salts thereof.

25. The method of Claim 1, wherein said active compound is selected from the group consisting of compounds of formula (XII), (XIII) and (XIV).

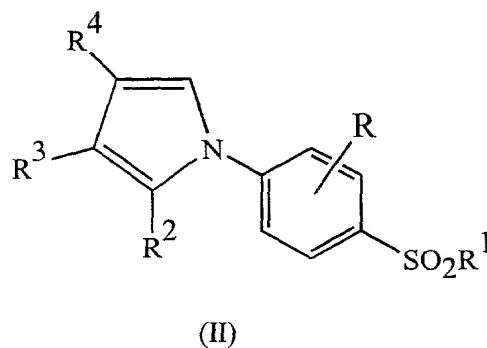
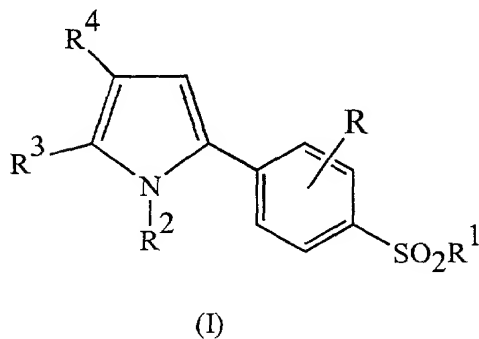
26. The method of Claim 25, wherein said active compound is selected from the group consisting of 3-phenyl-4-(4-methanesulfonylphenyl)-5H-furan-2-one and pharmaceutically acceptable salts thereof.

27. The method of Claim 25, wherein said active compound is selected from the group consisting of 2-(3,5-difluorophenyl)-3-(4-methanesulfonylphenyl)-2-cyclopenten-1-one and pharmaceutically acceptable salts thereof.

28. The method of Claim 25, wherein said active compound is selected from the

group consisting of 4-[5-methyl-3-phenylisoxazol-4-yl]benzenesulfonamide and pharmaceutically acceptable salts thereof.

29. A method for the treatment or prevention of tumor-related disorders in a mammal in need of such treatment or prevention, which method comprises
- 5 administering to said mammal an effective amount of an active compound selected from the group consisting of compounds of formula (I) and (II):



wherein

R represents a hydrogen atom, a halogen atom or a lower alkyl group;

- 10 R^1 represents a lower alkyl group, an amino group or a group of formula -
NHR^a (wherein R^a represents a group which may be eliminated *in vivo*);

R^2 represents a phenyl group or a phenyl group having at least one substituent selected from the group consisting of substituents α and substituents β , defined below;

- 15 R^3 represents a hydrogen atom, a halogen atom, a lower alkyl group or a lower alkyl group having at least one substituent selected from the group consisting of substituents α ;

- R^4 represents a hydrogen atom, a lower alkyl group, a lower alkyl group having at least one substituent selected from the group consisting of
- 20 substituents α , a cycloalkyl group, an aryl group as defined below, or an aralkyl group as defined below;

said aryl group is a carbocyclic aromatic hydrocarbon group having from 6 to 14 carbon atoms in one or more aromatic rings or such a group which is fused to a cycloalkyl group having from 3 to 10 carbon atoms, and the group is unsubstituted or it is substituted by at least one substituent selected from the group consisting of substituents α and substituents β ;

said aralkyl group is a lower alkyl group which is substituted by one or more of the aryl groups defined above;

said substituents α are selected from the group consisting of hydroxy groups, halogen atoms, lower alkoxy groups and lower alkylthio groups; and

said substituents β are selected from the group consisting of lower alkyl groups, alkanoyloxy groups, mercapto groups, alkanoylthio groups, lower alkylsulfinyl groups, lower alkyl groups having at least one substituent selected from the group consisting of substituents α , cycloalkyloxy groups, lower haloalkoxy groups and lower alkylenedioxy groups;

and pharmaceutically acceptable salts thereof.

30. The method of Claim 29, wherein R represents a hydrogen atom, a fluorine atom, a chlorine atom or a methyl group.

31. The method of Claim 29, wherein R represents a hydrogen atom.

32. The method of Claim 29, wherein R^1 represents a methyl group, an amino group or an acetylamino group.

33. The method of Claim 29, wherein R^1 represents an amino group or an acetylamino group.

34. The method of Claim 29, wherein R^2 represents a phenyl group or a phenyl group substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^1 .

substituents α^1 are selected from the group consisting of halogen atoms, lower

alkoxy groups and lower alkylthio groups; and

substituents β^1 are selected from the group consisting of lower alkyl groups, mercapto groups, alkanoylthio groups, lower alkyl groups substituted with at least one substituent selected from the group consisting of substituents α^1 ,
5 lower haloalkoxy groups and lower alkylenedioxy groups.

35. The method of Claim 29, wherein R^2 represents a phenyl group or a phenyl group substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^2 .

substituents α^1 are selected from the group consisting of halogen atoms, lower
10 alkoxy groups and lower alkylthio groups.

substituents β^2 are selected from the group consisting of lower alkyl groups, mercapto groups, alkanoylthio groups, lower alkyl groups substituted with a halogen atom, lower haloalkoxy groups and lower alkylenedioxy groups.

36. The method of Claim 29, wherein R^3 represents a hydrogen atom, a halogen
15 atom, a lower alkyl group or a lower alkyl group substituted with at least one substituent selected from the group consisting of substituents α^1 .

substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups.

37. The method of Claim 29, wherein R^3 represents a hydrogen atom, a halogen
20 atom, a lower alkyl group or a lower alkyl group substituted with a halogen atom.

38. The method of Claim 29, wherein R^4 represents a hydrogen atom, a lower alkyl group, a lower alkyl group substituted with at least one substituent selected from the group consisting of substituents α , a cycloalkyl group, an aryl group, an aryl group substituted with at least one substituent selected from the group consisting of
25 substituents α^1 and substituents β^3 , an aralkyl group or an aralkyl group substituted

with at least one substituent selected from the group consisting of substituents α^1 and substituents β^3 .

substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups; and

5 substituents β^3 are selected from the group consisting of lower alkyl groups, lower alkyl groups substituted with at least one substituent selected from the group consisting of substituents α and cycloalkyloxy groups.

39. The method of Claim 29, wherein R^4 represents a hydrogen atom, a lower alkyl group, a lower alkyl group substituted with at least one substituent selected from the
10 group consisting of substituents α^2 , a cycloalkyl group, an aryl group, an aryl group substituted with at least one substituent selected from the group consisting of substituents α^2 and substituents β^4 , an aralkyl group or an aralkyl group substituted with at least one substituent selected from the group consisting of substituents α^2 and substituents β^4 .

15 substituents α^2 are selected from the group consisting of hydroxy groups, halogen atoms and lower alkoxy groups; and

substituents β^4 are selected from the group consisting of lower alkyl groups, lower alkyl groups substituted with halogen atom and cycloalkyloxy groups.

40. The method of Claim 29, wherein said active compound is:

20 3-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,
4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,
1-(4-fluorophenyl)-2-(4-sulfamoylphenyl)pyrrole,
1-(4-fluorophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
5-fluoro-1-(4-fluorophenyl)-2-(4-methylsulfonylphenyl)pyrrole,

- 2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
1-(4-methoxyphenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
4-ethyl-2-(4-methoxyphenyl)-1-(4-sulfamoylphenyl)pyrrole,
2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
5 4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl)pyrrole,
2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
4-methyl-2-phenyl-1-(4-sulfamoylphenyl)pyrrole,
10 2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
2-(3-chloro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
4-methyl-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole,
5-chloro-1-(4-methoxyphenyl)-2-(4-sulfamoylphenyl)pyrrole,
1-(3,4-dimethylphenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
15 5-chloro-1-(4-ethoxyphenyl)-2-(4-sulfamoylphenyl)pyrrole,
5-chloro-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole,
1-(4-ethylthiophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
2-(3,5-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
1-(4-mercaptophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
20 1-(4-acetylthiophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(4-methoxyphenyl)pyrrole, or

1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(3,4-dimethylphenyl)pyrrole,

or a pharmaceutically acceptable salt thereof.

41. The method of Claim 29, wherein said active compound is:

2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

5 2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

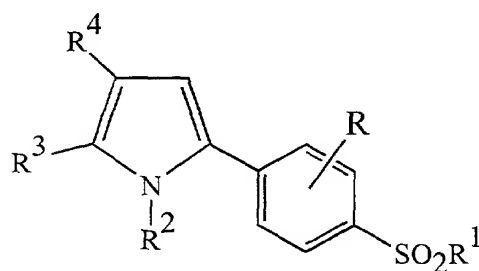
2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

4-methyl-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole, or

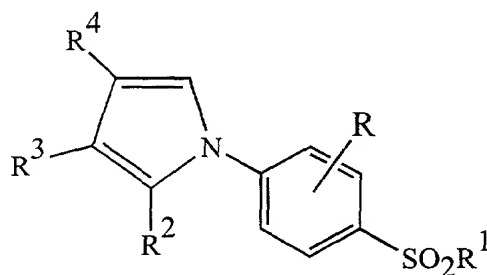
1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(3,4-dimethylphenyl)pyrrole,

or a pharmaceutically acceptable salt thereof.

10 42. A method for inhibiting tumor growth in a mammal in need thereof, which method comprises administering to said mammal an effective amount of an active compound selected from the group consisting of compounds of formula (I) and (II):



(I)



(II)

wherein

15 R represents a hydrogen atom, a halogen atom or a lower alkyl group;

R¹ represents a lower alkyl group, an amino group or a group of formula -NHR^a (wherein R^a represents a group which may be eliminated *in vivo*);

R² represents a phenyl group or a phenyl group having at least one substituent

selected from the group consisting of substituents α and substituents β , defined below;

R^3 represents a hydrogen atom, a halogen atom, a lower alkyl group or a lower alkyl group having at least one substituent selected from the group consisting of substituents α ;

R^4 represents a hydrogen atom, a lower alkyl group, a lower alkyl group having at least one substituent selected from the group consisting of substituents α , a cycloalkyl group, an aryl group as defined below, or an aralkyl group as defined below;

said aryl group is a carbocyclic aromatic hydrocarbon group having from 6 to 14 carbon atoms in one or more aromatic rings or such a group which is fused to a cycloalkyl group having from 3 to 10 carbon atoms, and the group is unsubstituted or it is substituted by at least one substituent selected from the group consisting of substituents α and substituents β ;

said aralkyl group is a lower alkyl group which is substituted by one or more of the aryl groups defined above;

said substituents α are selected from the group consisting of hydroxy groups, halogen atoms, lower alkoxy groups and lower alkylthio groups; and

said substituents β are selected from the group consisting of lower alkyl groups, alkanoyloxy groups, mercapto groups, alkanoylthio groups, lower alkylsulfinyl groups, lower alkyl groups having at least one substituent selected from the group consisting of substituents α , cycloalkyloxy groups, lower haloalkoxy groups and lower alkylenedioxy groups;

and pharmaceutically acceptable salts thereof.

43. The method of Claim 42, wherein R represents a hydrogen atom, a fluorine atom, a chlorine atom or a methyl group.

44. The method of Claim 42, wherein R represents a hydrogen atom.

45. The method of Claim 42, wherein R^1 represents a methyl group, an amino group or an acetylamino group.

46. The method of Claim 42, wherein R^1 represents an amino group or an acetylamino group.

5 47. The method of Claim 42, wherein R^2 represents a phenyl group or a phenyl group substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^1 .

substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups; and

10 substituents β^1 are selected from the group consisting of lower alkyl groups, mercapto groups, alkanoylthio groups, lower alkyl groups substituted with at least one substituent selected from the group consisting of substituents α^1 , lower haloalkoxy groups and lower alkylenedioxy groups.

15 48. The method of Claim 42, wherein R^2 represents a phenyl group or a phenyl group substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^2 .

substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups.

20 substituents β^2 are selected from the group consisting of lower alkyl groups, mercapto groups, alkanoylthio groups, lower alkyl groups substituted with a halogen atom, lower haloalkoxy groups and lower alkylenedioxy groups.

49. The method of Claim 42, wherein R^3 represents a hydrogen atom, a halogen atom, a lower alkyl group or a lower alkyl group substituted with at least one substituent selected from the group consisting of substituents α^1 .

25 substituents α^1 are selected from the group consisting of halogen atoms, lower

alkoxy groups and lower alkylthio groups.

50. The method of Claim 42, wherein R^3 represents a hydrogen atom, a halogen atom, a lower alkyl group or a lower alkyl group substituted with a halogen atom.

51. The method of Claim 42, wherein R^4 represents a hydrogen atom, a lower alkyl group, a lower alkyl group substituted with at least one substituent selected from the group consisting of substituents α , a cycloalkyl group, an aryl group, an aryl group substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^3 , an aralkyl group or an aralkyl group substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^3 .

substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups; and

substituents β^3 are selected from the group consisting of lower alkyl groups, lower alkyl groups substituted with at least one substituent selected from the group consisting of substituents α and cycloalkyloxy groups.

52. The method of Claim 42, wherein R^4 represents a hydrogen atom, a lower alkyl group, a lower alkyl group substituted with at least one substituent selected from the group consisting of substituents α^2 , a cycloalkyl group, an aryl group, an aryl group substituted with at least one substituent selected from the group consisting of substituents α^2 and substituents β^4 , an aralkyl group or an aralkyl group substituted with at least one substituent selected from the group consisting of substituents α^2 and substituents β^4 .

substituents α^2 are selected from the group consisting of hydroxy groups, halogen atoms and lower alkoxy groups; and

substituents β^4 are selected from the group consisting of lower alkyl groups, lower alkyl groups substituted with halogen atom and cycloalkyloxy groups.

53. The method of Claim 42, wherein said active compound is:

3-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,

4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,

1-(4-fluorophenyl)-2-(4-sulfamoylphenyl)pyrrole,

5 1-(4-fluorophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,

5-fluoro-1-(4-fluorophenyl)-2-(4-methylsulfonylphenyl)pyrrole,

2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

1-(4-methoxyphenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,

4-ethyl-2-(4-methoxyphenyl)-1-(4-sulfamoylphenyl)pyrrole,

10 2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl)pyrrole,

2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

15 4-methyl-2-phenyl-1-(4-sulfamoylphenyl)pyrrole,

2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(3-chloro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

4-methyl-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole,

5-chloro-1-(4-methoxyphenyl)-2-(4-sulfamoylphenyl)pyrrole,

20 1-(3,4-dimethylphenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,

5-chloro-1-(4-ethoxyphenyl)-2-(4-sulfamoylphenyl)pyrrole,

- 5-chloro-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole,
1-(4-ethylthiophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
2-(3,5-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
1-(4-mercaptophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
5 1-(4-acetylthiophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(4-methoxyphenyl)pyrrole, or
1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(3,4-dimethylphenyl)pyrrole,

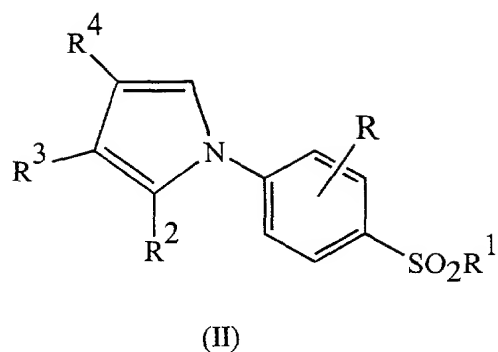
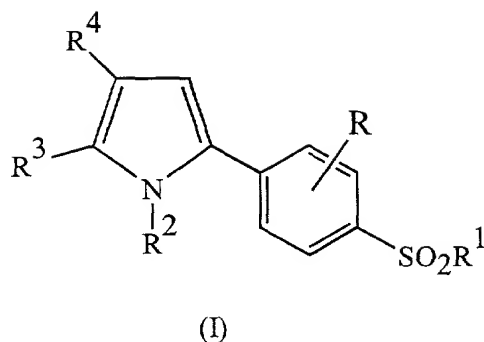
or a pharmaceutically acceptable salt thereof.

54. The method of Claim 42, wherein said active compound is:

- 10 2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
4-methyl-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole, or
1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(3,4-dimethylphenyl)pyrrole,

15 or a pharmaceutically acceptable salt thereof.

55. A method for inhibiting tumor metastasis in a mammal in need thereof, which method comprises administering to said mammal an effective amount of an active compound selected from the group consisting of compounds of formula (I) and (II):



wherein

R represents a hydrogen atom, a halogen atom or a lower alkyl group;

R¹ represents a lower alkyl group, an amino group or a group of formula -

5 NHR^a (wherein R^a represents a group which may be eliminated *in vivo*);

R² represents a phenyl group or a phenyl group having at least one substituent selected from the group consisting of substituents α and substituents β , defined below;

10 R³ represents a hydrogen atom, a halogen atom, a lower alkyl group or a lower alkyl group having at least one substituent selected from the group consisting of substituents α ;

15 R⁴ represents a hydrogen atom, a lower alkyl group, a lower alkyl group having at least one substituent selected from the group consisting of substituents α , a cycloalkyl group, an aryl group as defined below, or an aralkyl group as defined below;

20 said aryl group is a carbocyclic aromatic hydrocarbon group having from 6 to 14 carbon atoms in one or more aromatic rings or such a group which is fused to a cycloalkyl group having from 3 to 10 carbon atoms, and the group is unsubstituted or it is substituted by at least one substituent selected from the group consisting of substituents α and substituents β ;

said aralkyl group is a lower alkyl group which is substituted by one or more of

the aryl groups defined above;

said substituents α are selected from the group consisting of hydroxy groups, halogen atoms, lower alkoxy groups and lower alkylthio groups; and

5 said substituents β are selected from the group consisting of lower alkyl groups, alkanoyloxy groups, mercapto groups, alkanoylthio groups, lower alkylsulfinyl groups, lower alkyl groups having at least one substituent selected from the group consisting of substituents α , cycloalkyloxy groups, lower haloalkoxy groups and lower alkylenedioxy groups;

and pharmaceutically acceptable salts thereof.

10 56. The method of Claim 55, wherein R represents a hydrogen atom, a fluorine atom, a chlorine atom or a methyl group.

57. The method of Claim 55, wherein R represents a hydrogen atom.

58. The method of Claim 55, wherein R^1 represents a methyl group, an amino group or an acetylamino group.

15 59. The method of Claim 55, wherein R^1 represents an amino group or an acetylamino group.

60. The method of Claim 55, wherein R^2 represents a phenyl group or a phenyl group substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^1 .

20 substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups; and

substituents β^1 are selected from the group consisting of lower alkyl groups, mercapto groups, alkanoylthio groups, lower alkyl groups substituted with at least one substituent selected from the group consisting of substituents α^1 ,
25 lower haloalkoxy groups and lower alkylenedioxy groups.

61. The method of Claim 55, wherein R^2 represents a phenyl group or a phenyl group substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^2 .

5 substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups.

substituents β^2 are selected from the group consisting of lower alkyl groups, mercapto groups, alkanoylthio groups, lower alkyl groups substituted with a halogen atom, lower haloalkoxy groups and lower alkylenedioxy groups.

62. The method of Claim 55, wherein R^3 represents a hydrogen atom, a halogen atom, a lower alkyl group or a lower alkyl group substituted with at least one substituent selected from the group consisting of substituents α^1 .

substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups.

63. The method of Claim 55, wherein R^3 represents a hydrogen atom, a halogen atom, a lower alkyl group or a lower alkyl group substituted with a halogen atom.

64. The method of Claim 55, wherein R^4 represents a hydrogen atom, a lower alkyl group, a lower alkyl group substituted with at least one substituent selected from the group consisting of substituents α , a cycloalkyl group, an aryl group, an aryl group substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^3 , an aralkyl group or an aralkyl group substituted with at least one substituent selected from the group consisting of substituents α^1 and substituents β^3 .

substituents α^1 are selected from the group consisting of halogen atoms, lower alkoxy groups and lower alkylthio groups; and

25 substituents β^3 are selected from the group consisting of lower alkyl groups,

lower alkyl groups substituted with at least one substituent selected from the group consisting of substituents α and cycloalkyloxy groups.

65. The method of Claim 55, wherein R^4 represents a hydrogen atom, a lower alkyl group, a lower alkyl group substituted with at least one substituent selected from the group consisting of substituents α^2 , a cycloalkyl group, an aryl group, an aryl group substituted with at least one substituent selected from the group consisting of substituents α^2 and substituents β^4 , an aralkyl group or an aralkyl group substituted with at least one substituent selected from the group consisting of substituents α^2 and substituents β^4 .

10 substituents α^2 are selected from the group consisting of hydroxy groups, halogen atoms and lower alkoxy groups; and

 substituents β^4 are selected from the group consisting of lower alkyl groups, lower alkyl groups substituted with halogen atom and cycloalkyloxy groups.

66. The method of Claim 55, wherein said active compound is:

15 3-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,

 4-methyl-2-(4-methylphenyl)-1-(4-sulfamoylphenyl)pyrrole,

 1-(4-fluorophenyl)-2-(4-sulfamoylphenyl)pyrrole,

 1-(4-fluorophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,

 5-fluoro-1-(4-fluorophenyl)-2-(4-methylsulfonylphenyl)pyrrole,

20 2-(4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

 1-(4-methoxyphenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,

 4-ethyl-2-(4-methoxyphenyl)-1-(4-sulfamoylphenyl)pyrrole,

 2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

- 4-methyl-2-(4-methylthiophenyl)-1-(4-sulfamoylphenyl)pyrrole,
2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
2-(4-methoxy-3-methylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
2-(3-fluoro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
5 4-methyl-2-phenyl-1-(4-sulfamoylphenyl)pyrrole,
2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
2-(3-chloro-4-methoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
4-methyl-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole,
5-chloro-1-(4-methoxyphenyl)-2-(4-sulfamoylphenyl)pyrrole,
10 1-(3,4-dimethylphenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
5-chloro-1-(4-ethoxyphenyl)-2-(4-sulfamoylphenyl)pyrrole,
5-chloro-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole,
1-(4-ethylthiophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
2-(3,5-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,
15 1-(4-mercaptophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
1-(4-acetylthiophenyl)-4-methyl-2-(4-sulfamoylphenyl)pyrrole,
1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(4-methoxyphenyl)pyrrole, or
1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(3,4-dimethylphenyl)pyrrole,

or a pharmaceutically acceptable salt thereof.

- 20 67. The method of Claim 55, wherein said active compound is:

2-(4-chlorophenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(4-ethoxyphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

2-(3,4-dimethylphenyl)-4-methyl-1-(4-sulfamoylphenyl)pyrrole,

4-methyl-1-(4-methylthiophenyl)-2-(4-sulfamoylphenyl)pyrrole, or

1-(4-acetylaminosulfonylphenyl)-4-methyl-2-(3,4-dimethylphenyl)pyrrole,

5 or a pharmaceutically acceptable salt thereof.

ABSTRACT OF THE DISCLOSURE

Certain cyclooxygenase-2 inhibitors are useful for the treatment and prevention of tumors and tumor-related disorders and cachexia.

1. A method of treating or preventing a tumor or tumor-related disorder or cachexia, comprising administering to a subject in need thereof a therapeutically effective amount of a cyclooxygenase-2 inhibitor.

APPLICATION FOR UNITED STATES LETTERS PATENT**Declaration and Power of Attorney**

As a below named inventor, I declare that:

My residence, post office address and citizenship are as stated below next to my name; I believe that I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

Methods and compositions for the treatment and prevention of tumors,
the specification of which is attached hereto. tumor-related disorders and cachexia

I have reviewed and understand the contents of said specification, including the claims.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR §1.56.

I claim priority benefits under 35 USC §119 of: (i) any foreign application(s) for patent or inventor's certificate listed below; or (ii) any United States provisional application(s) listed below; and have also identified below any foreign application(s) for patent or inventor's certificate, or PCT international application having a filing date before that of the application(s) on which priority is claimed.

COUNTRY	APPLICATION NUMBER	DATE (day, month, year)	PRIORITY CLAIMED
Japan	Hei 9-354499	24 Dec. 1997	yes <input checked="" type="checkbox"/> no <input type="checkbox"/>
Japan	Hei 10-15306	28 Jan. 1998	yes <input checked="" type="checkbox"/>
Japan	Hei 10-204907	21 Jul. 1998	yes <input checked="" type="checkbox"/> no <input type="checkbox"/>
Japan	Hei 10-269444	24 Sep. 1998	yes <input checked="" type="checkbox"/>

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.


I appoint the following attorneys to prosecute this application and to transact all business in the U.S. Patent & Trademark Office connected therewith: Stephen H. Frishauf, Reg. No. 16,233; Leonard Holtz, Reg. No. 22,974; Herbert Goodman, Reg. No. 17,081; Thomas Langer, Reg. No. 27,264; Marshall J. Chick, Reg. No. 26,853; Walter J. Baum, Reg. No. 20,641; Richard S. Barth, Reg. No. 28,180; Douglas Holtz, Reg. No. 33,902; and Robert P. Michal, Reg. No. 35,614.

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